

NEW SUBSTITUTED INDOLINONES, PREPARATION THEREOF
AND THEIR USE AS PHARMACEUTICAL COMPOSITIONS

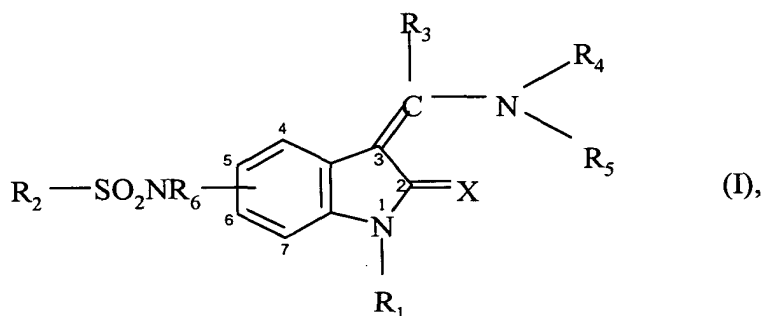
Related Applications

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This is a division of Serial No. 10/002,939 filed November 1, 2001. Benefit of U.S. Provisional Application Serial No. 60/251,055, filed on December 1, 2000 is hereby claimed.

10 Description of the Invention

The present invention relates to new substituted indolinones of general formula



15 the isomers, the salts thereof, particularly the physiologically acceptable salts thereof which have valuable properties.

The above compounds of general formula I wherein R_1 is a hydrogen atom or a prodrug group have valuable pharmacological properties, particularly an inhibiting effect on the proliferation of cultivated human tumour cells, but also on the proliferation of other cells, particularly endothelial cells, e.g. in angiogenesis, on various kinases, particularly on
20 receptor tyrosine kinases (such as, for example, VEGFR2, EGFR, IGF1R), non-receptor tyrosine kinases (such as e.g. c-src), and serine/threonine kinases (such as e.g. cyclin-dependent kinases), and the other compounds of the above general formula I wherein R_1 does not denote a hydrogen atom or a prodrug group, are valuable intermediate products
25 for the preparation of the compounds mentioned above.

The present invention thus relates to the above compounds of general formula I, wherein

- X denotes an oxygen or sulphur atom,
- R₁ denotes a hydrogen atom, a C₁₋₄-alkoxycarbonyl or C₂₋₄-alkanoyl group,
- R₂ denotes a C₁₋₆-alkyl group optionally substituted by one or more halogen atoms or
5 a phenyl group or a C₂₋₆-alkenyl group optionally substituted by a phenyl group, wherein
the phenyl moiety may be substituted in each case by a fluorine, chlorine, bromine or
iodine atom, by a C₁₋₃-alkyl or C₁₋₃-alkoxy group,
a phenyl group which may be mono- or disubstituted by fluorine, chlorine, bromine or
iodine atoms, by C₁₋₃-alkyl or C₁₋₃-alkoxy groups, wherein the substituents may be
10 identical or different,
a phenyl group substituted by a trifluoromethyl, carboxy, C₁₋₃-alkoxycarbonyl,
aminocarbonyl, cyano, aminomethyl, nitro or amino group,
a C₄₋₆-alkyl, C₃₋₇-cycloalkyl, trimethylphenyl or naphthyl group,
a 5-membered heteroaromatic group optionally substituted by a C₁₋₃-alkyl group, which
15 contains, in the heteroaromatic moiety,
an imino group optionally substituted by a C₁₋₃-alkyl group, an oxygen or sulphur atom,
an imino group optionally substituted by a C₁₋₃-alkyl group and an oxygen, sulphur or
nitrogen atom,
an imino group optionally substituted by a C₁₋₃-alkyl group and two nitrogen atoms, or
20 an oxygen or sulphur atom and two nitrogen atoms, and to which a phenyl ring may be
fused via two adjacent carbon atoms,
or denotes a 6-membered heteroaromatic group optionally substituted by a C₁₋₃-alkyl
group, which contains one or two heteroatoms in the heteroaromatic moiety and to which
a phenyl ring may be fused via two adjacent carbon atoms,
- 25 R₃ denotes a hydrogen atom or a C₁₋₆-alkyl group,
a phenyl group optionally substituted by a fluorine, chlorine or bromine atom, by a C₁₋₃-
alkyl, hydroxy, C₁₋₃-alkoxy, C₁₋₃-alkylsulphenyl, C₁₋₃-alkylsulphinyl, C₁₋₃-alkylsulphonyl,
phenylsulphenyl, phenylsulphinyl, phenylsulphonyl, nitro, amino, C₁₋₃-alkylamino, di-
(C₁₋₃-alkyl)-amino, C₂₋₅-alkanoylamino or N-(C₁₋₃-alkylamino)-C₂₋₅-alkanoylamino
30 group,
- R₄ denotes a phenyl or naphthyl group optionally substituted by R₇, which may
additionally be substituted by a chlorine or bromine atom or a nitro group, a 5-membered

heteroaromatic group which contains an imino group, an oxygen or sulphur atom or an imino group, an oxygen or sulphur atom and one or two nitrogen atoms, or a 6-membered heteroaromatic group which contains one, two or three nitrogen atoms, while the abovementioned 5- and 6-membered heteroaromatic groups may additionally be substituted by a chlorine or bromine atom or by a methyl group or wherein a phenyl ring may be fused to the abovementioned 5- and 6-membered heteroaromatic groups via 2 adjacent carbon atoms, or

R₅ and R₆ in each case independently of one another denote hydrogen atoms or C₁₋₃-alkyl groups, and

R₇ denotes a fluorine, chlorine, bromine or iodine atom or a cyano group, a methoxy group or a C₂₋₃-alkoxy group, which may be substituted in the 2 or 3 position by an amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino or 5- to 7-membered cycloalkyleneimino group, while in each case an alkyl moiety in the abovementioned alkylamino and dialkylamino groups may additionally be substituted by a phenyl group, a trifluoromethyl, nitro, amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, C₂₋₅-alkanoylamino, N-(C₁₋₃-alkyl)-C₂₋₅-alkanoylamino, C₁₋₅-alkylsulphonylamino, N-(C₁₋₃-alkyl)-C₁₋₅-alkylsulphonylamino, phenylsulphonylamino, N-(C₁₋₃-alkyl)-phenylsulphonylamino, aminosulphonyl, C₁₋₃-alkylaminosulphonyl or di-(C₁₋₃-alkyl)-aminosulphonyl group, while in each case an alkyl moiety in the abovementioned alkylamino and dialkylamino groups may additionally be substituted by a carboxy, C₁₋₃-alkoxycarbonyl, aminocarbonyl, C₁₋₃-alkylaminocarbonyl, di-(C₁₋₃-alkyl)-aminocarbonyl, 2-dimethylaminoethylaminocarbonyl or N-methyl-(2-dimethylaminoethyl)-aminocarbonyl group and in each case the alkyl moiety of the abovementioned alkanoylamino or alkylsulphonylamino groups may additionally be substituted by a phenyl, amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino or a 4- to 7-membered cycloalkyleneimino group, a C₂₋₄-alkylamino group which is terminally substituted in the 2, 3- or 4 position by an amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, benzylamino, N-(C₁₋₃-alkyl)-benzylamino, C₂₋₅-alkanoylamino or N-(C₁₋₃-alkyl)-C₂₋₅-alkanoylamino group and wherein additionally the amino-hydrogen atom may be replaced by a C₂₋₅-alkanoyl, benzoyl, C₁₋₅-alkylsulphonyl- or phenylsulphonyl group, while the last-mentioned C₂₋₅-alkanoyl or C₁₋₅-alkylsulphonyl groups in the alkyl moiety may be substituted by a phenyl group,

a carbonyl group which is substituted by a hydroxy, C₁₋₃-alkoxy, amino, C₁₋₃-alkylamino, N-(C₁₋₅-alkyl)-C₁₋₃-alkylamino or C₅₋₇-cycloalkyleneimino group;

a C₁₋₃-alkyl group which may be substituted by an amino, C₁₋₅-alkylamino,

C₅₋₇-cycloalkylamino or phenyl-C₁₋₃-alkylamino group which may additionally be

5 substituted at the amino nitrogen atom in each case by a C₁₋₄-alkyl, C₅₋₇-cycloalkyl or C₂₋₄-alkenyl- or C₁₋₄-alkyl group, while

the abovementioned C₁₋₄-alkyl substituent in each case may additionally be mono-, di- or trisubstituted by a cyano, carboxy, C₁₋₃-alkoxycarbonyl, C₂₋₄-alkanoyl, pyridyl,

imidazolyl, benzo[1,3]dioxol or phenyl group, while the phenyl group may be substituted

10 by fluorine, chlorine or bromine atoms, by methyl, methoxy, trifluoromethyl, cyano or nitro groups and the substituents may be identical or different, or in the 2, 3 or 4 position by a hydroxy group,

a C₁₋₃-alkyl group which is substituted by a hydroxy, carboxy, morpholino,

thiomorpholino, 1-oxo-thiomorpholino, 1,1-dioxo-thiomorpholino, piperazino, N-(C₁₋₃-

15 alkyl)-piperazino or N-benzyl-piperazino group, by a 5- to 7-membered

cycloalkenyleneimino group or by a 4- to 7-membered cycloalkyleneimino group, while

the abovementioned 5- to 7-membered cycloalkyleneimino groups may be substituted by one or two C₁₋₃-alkyl groups, which may in turn be terminally substituted by a hydroxy,

amino or C₂₋₄-alkanoylamino group, or by a C₅₋₇-cycloalkyl or phenyl group and by a

20 hydroxy group and in the abovementioned cycloalkyleneimino groups a methylene group adjacent to the nitrogen atom may be replaced by a carbonyl group,

a C₁₋₃-alkyl group which is substituted by a 5- to 7-membered cycloalkyleneimino group,

while a phenyl group optionally mono- or disubstituted by fluorine, chlorine or bromine atoms or by methyl or methoxy groups, wherein the substituents may be identical or

25 different, or an oxazolo, imidazolo, thiazolo, pyridino, pyrazino or pyrimidino group optionally substituted by a fluorine, chlorine, bromine or iodine atom, by a methyl,

methoxy or amino group is fused to the abovementioned 5- to 7-membered

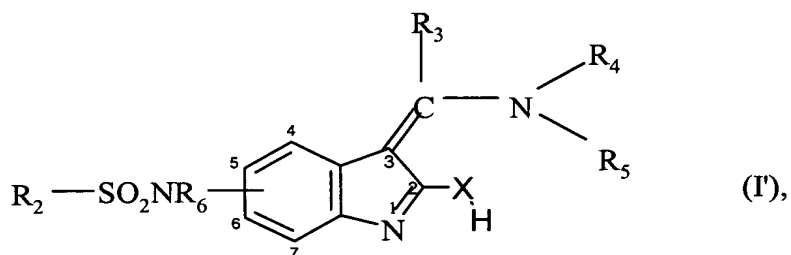
cycloalkyleneimino groups via 2 adjacent carbon atoms, while the abovementioned

monosubstituted phenyl groups may additionally be substituted by a fluorine, chlorine or

30 bromine atom, by a methyl, methoxy or nitro group, or

denotes an imidazolyl or 1H-C₁₋₃-alkylimidazolyl group.

If R_1 denotes a hydrogen atom, the present invention also relates to the tautomeric compounds of formula I'



5 The invention also relates to compounds of formula I, wherein R_1 denotes a cleavable prodrug group.

The invention further relates to pharmaceutical compositions containing the pharmacologically active compound, their use and processes for preparing them.

10 Preferred compounds of formula I are those wherein the sulphonylamino group of formula R_2 -SO₂NR₆- is linked to the 5-position of the indolinone group.

Preferred compounds of formula I are those wherein

R_7 denotes a C₁₋₃-alkyl group which is substituted by a hydroxy, carboxy, morpholino, thiomorpholino, 1-oxo-thiomorpholino, 1,1-dioxo-thiomorpholino, piperazino, N-(C₁₋₃-alkyl)-piperazino or N-benzyl-piperazino group, by a 5- to 7-membered cycloalkenyleneimino group or by a 4- to 7-membered cycloalkyleneimino group, while the abovementioned 5- to 7-membered cycloalkyleneimino groups may be substituted by one or two C₁₋₃-alkyl groups, which may in turn be terminally substituted by an amino or C₂₋₄-alkanoylamino group, or by a C₅₋₇-cycloalkyl or phenyl group and by a hydroxy group and in the abovementioned cycloalkyleneimino groups a methylene group adjacent to the nitrogen atom may be replaced by a carbonyl group.

Also preferred are compounds of formula I wherein

25 R_3 denotes a phenyl group optionally substituted by a fluorine, chlorine or bromine atom, by a C₁₋₃-alkyl, hydroxy, C₁₋₃-alkoxy, C₁₋₃-alkylsulphenyl, C₁₋₃-alkylsulphinyl, C₁₋₃-alkylsulphonyl, phenylsulphenyl, phenylsulphinyl, phenylsulphonyl, nitro, amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, C₂₋₅-alkanoylamino or N-(C₁₋₃-alkylamino)-C₂₋₅-

alkanoylamino group, more particularly a phenyl group optionally substituted by an fluorine, chlorine, bromine or iodine atom, by a C₁₋₃-alkyl, C₁₋₃-alkoxy, nitro or amino group.

- 5 In another preferred embodiment R₂ denotes a C₁₋₄-alkyl group optionally substituted by one or more halogen atoms or a phenyl group, a C₃₋₅-cycloalkyl group or a C₂₋₄-alkenyl group optionally substituted by a phenyl group, wherein the phenyl moiety in each case may be substituted by a fluorine, chlorine, bromine or iodine atom or by a C₁₋₃-alkyl or C₁₋₃-alkoxy.

10

Moreover, the carboxy, amino or imino groups present in a compound of the above general formula I may be substituted by groups which can be cleaved *in vivo*.

- In addition to the alkoxycarbonyl and alkanoyl groups already mentioned hereinbefore, groups which can be cleaved *in vivo* may also be included, such as an acyl group such as the benzoyl, pyridinoyl, pentanoyl or hexanoyl group, an allyloxycarbonyl group, a C₁₋₁₆-alkoxycarbonyl group such as the *tert.*-butyloxycarbonyl, pentyloxycarbonyl, hexyloxycarbonyl, octyloxycarbonyl, nonyloxycarbonyl, decyloxycarbonyl, undecyloxycarbonyl, dodecyloxycarbonyl or hexadecyloxycarbonyl group, a phenyl-C₁₋₆-alkoxycarbonyl group such as the benzyloxycarbonyl, phenylethoxycarbonyl or phenylpropoxycarbonyl group, a C₁₋₃-alkylsulphonyl-C₂₋₄-alkoxycarbonyl, C₁₋₃-alkoxy-C₂₋₄-alkoxy-C₂₋₄-alkoxycarbonyl or R_cCO-O-(R_dCR_e)-O-CO-group, wherein
- 15
- 20

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R_c denotes a C₁₋₈-alkyl, C₅₋₇-cycloalkyl, phenyl- or phenyl-C₁₋₃-alkyl group,

R_e denotes a hydrogen atom, a C₁₋₃-alkyl, C₅₋₇-cycloalkyl or phenyl group and

R_d denotes a hydrogen atom or a C₁₋₃-alkyl group or a R_fCO-O-(R_gCR_h)-O-Rest wherein

R_f denotes a C₁₋₈-alkyl, C₅₋₇-cycloalkyl, phenyl or phenyl-C₁₋₃-alkyl group,

30

R_g denotes a hydrogen atom, a C₁₋₃-alkyl, C₅₋₇-cycloalkyl or phenyl group and

R_h denotes a hydrogen atom or a C₁₋₃-alkyl group,

while the abovementioned ester groups may also be used as a group which can be converted *in vivo* into a carboxy group.

5

Preferred compounds of the above general formula I are those wherein

X denotes an oxygen atom,

R₁ denotes a hydrogen atom,

10 R₂ denotes a C₁₋₃-alkyl group optionally substituted by one or more fluorine atoms or a phenyl group or a C₂₋₄-alkenyl group optionally substituted by a phenyl group; a phenyl group which may be mono- or disubstituted by fluorine, chlorine, bromine or iodine atoms, by C₁₋₃-alkyl or C₁₋₃-alkoxy groups, wherein the substituents may be identical or different,

a phenyl group substituted by a trifluoromethyl, carboxy, C₁₋₃-alkoxycarbonyl,

15 aminocarbonyl, cyano, aminomethyl, nitro or amino group,

a C₄₋₆-alkyl, C₃₋₇-cycloalkyl, trimethylphenyl or naphthyl group, or

a pyridinyl, quinolyl, isoquinolyl, oxazolyl, isoxazolyl, imidazolyl or 1-(C₁₋₃-alkyl)-imidazolyl group optionally substituted by a C₁₋₃-alkyl group,

R₃ denotes a hydrogen atom or a C₁₋₄-alkyl group, or

20 a phenyl group optionally substituted by a fluorine, chlorine, bromine or iodine atom, by a C₁₋₃-alkyl, C₁₋₃-alkoxy, nitro or amino group,

R₄ denotes a phenyl group optionally substituted by R₇, which may additionally be substituted by a chloro or nitro group,

R₅ and R₆ in each case denote a hydrogen atom, and

25 R₇ denotes a fluorine, chlorine, bromine or iodine atom,

a methoxy, nitro, cyano, carboxy, C₁₋₃-alkoxycarbonyl, aminocarbonyl,

C₁₋₃-alkylaminocarbonyl, di-(C₁₋₃-alkyl)-aminocarbonyl, phenyl-C₁₋₃-alkylaminocarbonyl, N-(phenyl-C₁₋₃-alkyl)-C₁₋₃-alkylaminocarbonyl or 5- to 7-membered cycloalkyleneiminocarbonyl group,

30 a C₁₋₃-alkyl group which is substituted by a carboxy, C₁₋₃-alkoxycarbonyl,

aminocarbonyl, C₁₋₃-alkylaminocarbonyl, di-(C₁₋₃-alkyl)-aminocarbonyl, phenyl-C₁₋₃-alkylaminocarbonyl, N-(phenyl-C₁₋₃-alkyl)-C₁₋₃-alkylaminocarbonyl, 5- to 7-membered

cycloalkyleneiminocarbonyl, amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, phenyl-C₁₋₃-alkylamino, N-(phenyl-C₁₋₃-alkyl)-C₁₋₃-alkylamino or 5- to 7-membered cycloalkyleneimino group,

while the abovementioned 5- to 7-membered cycloalkyleneimino group may be substituted by one or two C₁₋₃-alkyl groups, which may in turn be terminally substituted by a hydroxy, amino or C₂₋₄-alkanoylamino group, and at the same time in the abovementioned 5- to 7-membered cycloalkyleneimino moieties a methylene group in the 2 position may be replaced by a carbonyl group or in the abovementioned 6- and 7-membered cycloalkyleneimino moieties a methylene group in the 4 position may be replaced by an oxygen atom, by an imino, N-(C₁₋₃-alkyl)-imino, N-(phenyl-C₁₋₃-alkyl)-imino or N-(C₁₋₅-alkoxycarbonyl)-imino group,

an amino, C₁₋₃-alkylamino, phenyl-C₁₋₃-alkylamino, C₁₋₅-alkanoylamino, phenyl-C₁₋₄-alkanoylamino, C₁₋₅-alkoxycarbonylamino, phenyl-C₁₋₃-alkoxycarbonylamino, C₁₋₅-alkylsulphonylamino, phenyl-C₁₋₃-alkylsulphonylamino- or phenylsulphonylamino group, wherein the hydrogen atom of the amino group may be replaced by a C₁₋₃-alkyl group, while the C₁₋₃-alkyl moiety may be substituted by a carboxy, C₁₋₃-alkoxycarbonyl, aminocarbonyl, C₁₋₃-alkylaminocarbonyl, di-(C₁₋₃-alkyl)-aminocarbonyl, phenyl-C₁₋₃-alkylaminocarbonyl, N-(phenyl-C₁₋₃-alkyl)-C₁₋₃-alkylaminocarbonyl, 2-dimethylaminoethylaminocarbonyl, N-methyl-(2-dimethylaminoethyl)-aminocarbonyl- or C₄₋₆-cycloalkyleneiminocarbonyl group or from position 2 by an amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, phenyl-C₁₋₃-alkylamino, N-(phenyl-C₁₋₃-alkyl)-C₁₋₃-alkylamino, C₂₋₅-alkanoylamino, N-(C₁₋₃-alkyl)-C₂₋₅-alkanoylamino, C₁₋₅-alkoxycarbonylamino- or N-(C₁₋₅-alkoxycarbonyl)-C₁₋₃-alkylamino group, imidazolyl or 1-C₁₋₃-alkylimidazolyl group.

Particularly preferred compounds of formula I are those wherein

R₇ denotes a C₁₋₃-alkyl group which is substituted by a carboxy, C₁₋₃-alkoxycarbonyl, aminocarbonyl, C₁₋₃-alkylaminocarbonyl, di-(C₁₋₃-alkyl)-aminocarbonyl, phenyl-C₁₋₃-alkylaminocarbonyl, N-(phenyl-C₁₋₃-alkyl)-C₁₋₃-alkylaminocarbonyl, 5- to 7-membered cycloalkyleneiminocarbonyl, amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, phenyl-C₁₋₃-alkylamino, N-(phenyl-C₁₋₃-alkyl)-C₁₋₃-alkylamino- or 5- to 7-membered cycloalkyleneimino group,

while the abovementioned 5- to 7-membered cycloalkyleneimino group may be substituted by one or two C₁₋₃-alkyl groups, which may in turn be terminally substituted by an amino or C₂₋₄-alkanoylamino group, and at the same time in the abovementioned 5- to 7-membered cycloalkyleneimino moieties a methylene group may be replaced in the 2 position by a carbonyl group or in the abovementioned 6- and 7-membered cycloalkyleneimino moieties a methylene group in the 4 position may be replaced by an oxygen atom, by an imino, N-(C₁₋₃-alkyl)-imino, N-(phenyl-C₁₋₃-alkyl)-imino or N-(C₁₋₅-alkoxycarbonyl)-imino group.

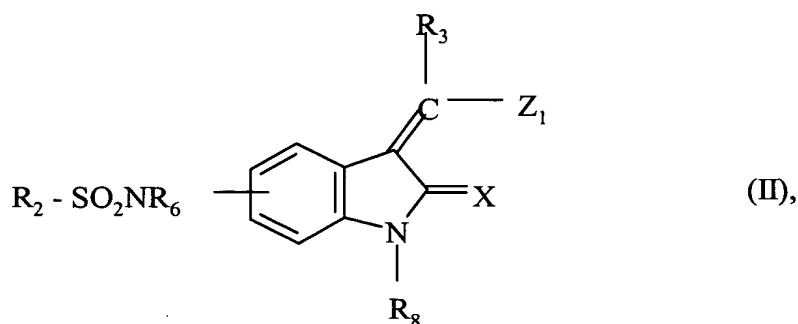
- 10 Particularly preferred compounds of general formula I are those wherein
- X denotes an oxygen atom,
- R₁ denotes a hydrogen atom,
- R₂ denotes a C₁₋₃-alkyl group optionally substituted by a phenyl group, a C₁₋₃-perfluoroalkyl group or a phenylvinyl group;
- 15 a phenyl group which may be substituted by a fluorine, chlorine, bromine or iodine atom, by a C₁₋₃-alkyl, C₁₋₃-alkoxy, nitro, amino, cyano or aminomethyl group, a C₄₋₆-alkyl, C₃₋₇-cycloalkyl, trimethylphenyl or naphthyl group, a pyridinyl, quinolyl, isoquinolyl, oxazolyl, isoxazolyl, imidazolyl or 1-(C₁₋₃-alkyl)-imidazolyl group optionally substituted by a C₁₋₃-alkyl group,
- 20 R₃ denotes a phenyl group optionally substituted by a fluorine, chlorine, bromine or iodine atom, by a C₁₋₃-alkyl, C₁₋₃-alkoxy, nitro or amino group,
- R₄ denotes a phenyl group which may be substituted by R₇ and additionally by a chlorine atom or a nitro group, while
- R₇ denotes a fluorine, chlorine, bromine or iodine atom,
- 25 a methoxy, nitro, cyano, carboxy, methoxycarbonyl, aminocarbonyl, methylaminocarbonyl, dimethylaminocarbonyl, benzylaminocarbonyl, N-benzyl-methylaminocarbonyl, pyrrolidinocarbonyl or piperidinocarbonyl group, a methyl or ethyl group which may be substituted by a carboxy, methoxycarbonyl, aminocarbonyl, methylaminocarbonyl, dimethylaminocarbonyl, benzylaminocarbonyl,
- 30 N-benzyl-methylaminocarbonyl, pyrrolidinocarbonyl, piperidinocarbonyl, amino, C₁₋₄-alkylamino, di-C₁₋₄-alkylamino, benzylamino, N-benzyl-C₁₋₄-alkylamino, C₂₋₄-alkanoylamino, N-C₁₋₄-alkyl-C₂₋₄-alkanoylamino, tert.butylloxycarbonylamino, N-

methyl-tert.butyloxycarbonylamino, pyrrolidino, pyrrolidinomethyl,
hydroxypyrrolidinomethyl, hydroxymethylpyrrolidinomethyl, piperidino, 4-(3-
aminopropyl)-piperidino, 4-(3-acetylaminopropyl)-piperidino, dimethylpiperidino, 2-oxo-
piperidino, piperazino, 4-methyl-piperazino, 4-benzyl-piperazino, 4-tert.butoxycarbonyl-
5 piperazino or morpholino group, or
an amino, methylamino, ethylamino, C₁₋₃-alkanoylamino, phenylacetylamino,
tert.butoxycarbonylamino, piperidinomethylcarbonylamino, C₁₋₄-alkylsulphonylamino,
phenyl-methylsulphonylamino or phenylsulphonylamino group, wherein the hydrogen
atom of the amino group may be replaced by a methyl, ethyl or propyl group, while the
10 methyl or ethyl moiety in each case may be substituted by a carboxy, methoxycarbonyl,
aminocarbonyl, methylaminocarbonyl, dimethylaminocarbonyl,
2-dimethylaminoethylaminocarbonyl or N-methyl-(2-dimethylaminoethyl)-
aminocarbonyl group or the ethyl moiety may also be substituted from position 2 by an
amino, methylamino, dimethylamino, benzylalkylamino, N-benzyl-methylamino, C₂₋₃-
15 alkanoylamino, N-methyl-C₂₋₃-alkanoylamino, tert.butyloxycarbonylamino or N-methyl-
tert.butyloxycarbonylamino group,
an imidazolyl or 1-methylimidazolyl group,
R₅ and R₆ in each case denote a hydrogen atom,
20 and the isomers and the salts thereof.

Particularly preferred are compounds of formula I wherein R₄ denotes a phenyl group
substituted by R₇ in the 3 or 4 position, particularly in the 4 position.

25 According to the invention, the new compounds are obtained, for example, by the
following methods known in principle from the literature:

a. reacting a compound of general formula



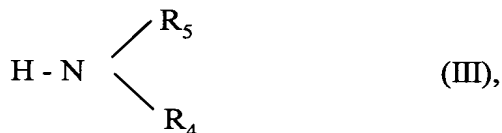
wherein

X, R₂, R₃ and R₆ are as hereinbefore defined and

R₈ has one of the meanings given for R₁ or may denote a protecting group for the
 5 nitrogen atom of the lactam group, while R₈ may also represent a bond to a solid phase optionally formed via a spacer, and

Z₁ denotes a halogen atom, a hydroxy, alkoxy or aralkoxy group, e.g. a chlorine or bromine atom, a methoxy, ethoxy or benzyloxy group,

10 with an amine of general formula



wherein

R₄ and R₅ are as hereinbefore defined,

and if necessary subsequently cleaving any protecting group used for the nitrogen atom of
 15 the lactam group or from a solid phase.

The protecting group used for the nitrogen atom of the lactam group may be, for example, an acetyl, benzoyl, ethoxycarbonyl, tert.butyloxycarbonyl or benzyloxycarbonyl group and

20 the solid phase used may be a resin such as a 4-(2',4'-dimethoxyphenylaminomethyl)-phenoxy resin, while the bond may expediently be effected via the amino group, or a p-benzyloxybenzyl alcohol resin, while the bond may expediently be effected via an intermediate member such as a 2,5-dimethoxy-4-hydroxy-benzyl derivative.

The reaction is conveniently carried out in a solvent such as dimethylformamide, toluene, acetonitrile, tetrahydrofuran, dimethylsulphoxide, dichloromethane or mixtures thereof, optionally in the presence of an inert base such as triethylamine, N-ethyl-diisopropylamine or sodium hydrogen carbonate at temperatures between 20 and 175°C, while any protecting group used may simultaneously be cleaved by transamidation.

If Z_1 in a compound of general formula II denotes a halogen atom, the reaction is preferably carried out in the presence of an inert base at temperatures between 20 and 120°C.

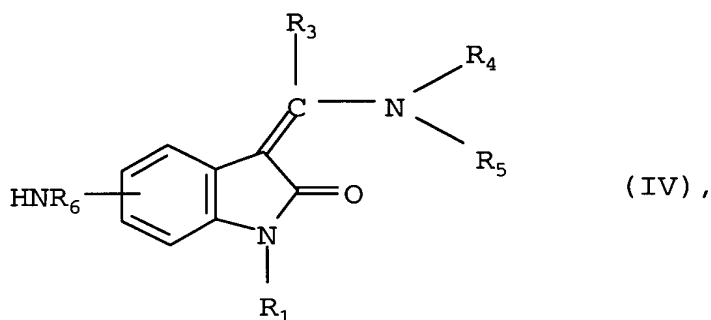
If Z_1 in a compound of general formula II denotes a hydroxy, alkoxy or aralkoxy group, the reaction is preferably carried out at temperatures between 20 and 200°C.

If any protecting group used subsequently has to be cleaved, this is conveniently carried out either hydrolytically in an aqueous or alcoholic solvent, e.g. in methanol/water, ethanol/water, isopropanol/water, tetrahydrofuran/water, dioxane/water, dimethylformamide/water, methanol or ethanol in the presence of an alkali metal base such as lithium hydroxide, sodium hydroxide or potassium hydroxide at temperatures between 0 and 100°C, preferably at temperatures between 10 and 50°C,

or advantageously by transamidation with an organic base such as ammonia, methylamine, butylamine, dimethylamine or piperidine in a solvent such as methanol, ethanol, dimethylformamide and mixtures thereof or in an excess of the amine used at temperatures between 0 and 100°C, preferably at temperatures between 10 and 50°C.

Any solid phase used is preferably cleaved using trifluoroacetic acid and water at temperatures between 0 and 35°C, preferably at ambient temperature.

b. reacting a compound of general formula



wherein

R_1 and R_3 to R_6 are as hereinbefore defined, with a compound of general formula

$R_2 - SO_2 - OH$ (V),

5 wherein

R_2 is as hereinbefore defined, or with the reactive derivatives thereof.

The reaction is preferably carried out in a solvent such as dichloromethane, diethylether, tetrahydrofuran, toluene, dioxane, acetonitrile, dimethylsulphoxide or

10 dimethylformamide, optionally with a reactive derivative of a compound of general formula V such as the halide thereof, in the presence of an inorganic or tertiary organic base, preferably at temperatures between 0°C and the boiling temperature of the solvent used, preferably at temperatures between 50 and 100°C.

15 With a corresponding sulphonic acid the reaction is preferably carried out in the presence of a dehydrating agent, e.g. in the presence of isobutyl chloroformate, tetraethyl orthocarbonate, trimethyl orthoacetate, 2,2-dimethoxypropane, tetramethoxysilane, thionyl chloride, trimethylchlorosilane, phosphorus trichloride, phosphorus pentoxide, N,N'-dicyclohexylcarbodiimide, N,N'-dicyclohexylcarbodiimide/N-hydroxysuccinimide, 20 N,N'-dicyclohexylcarbodiimide/1-hydroxy-benzotriazole, 2-(1H-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium-tetrafluoroborate, 2-(1H-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium-tetrafluoroborate/1-hydroxy-benzotriazole, N,N'-carbonyldiimidazole or triphenylphosphine/carbon tetrachloride, and optionally with the addition of a base such as pyridine, 4-dimethylaminopyridine, N-methyl-morpholine or 25 triethylamine, expediently at temperatures between 0 and 150°C, preferably at temperatures between 0 and 100°C.

If according to the invention a compound of general formula I is obtained which contains an alkoxy carbonyl group, this can be converted by hydrolysis into a corresponding carboxy compound, or

- 5 if a compound of general formula I is obtained which contains an amino or alkylamino group, this may be converted by reductive alkylation into a corresponding alkylamino or dialkylamino compound, or

- if a compound of general formula I is obtained which contains an amino or alkylamino
10 group, this may be converted by acylation into a corresponding acyl compound, or

- if a compound of general formula I is obtained which contains a carboxy group, this may be converted by esterification or amidation into a corresponding ester or aminocarbonyl compound, or

- 15 if a compound of general formula I is obtained which contains a nitro group, this can be converted by reduction into a corresponding amino compound, or

- if a compound of general formula I is obtained which contains a cyano group, this can be
20 converted by reduction into a corresponding aminomethyl compound.

- The subsequent hydrolysis is preferably carried out in an aqueous solvent, e.g. in water, isopropanol/water, tetrahydrofuran/water or dioxane/water, in the presence of an acid such as trifluoroacetic acid, hydrochloric acid or sulphuric acid or in the presence of an
25 alkali metal base such as lithium hydroxide, sodium hydroxide or potassium hydroxide at temperatures between 0 and 100°C, preferably at temperatures between 10 and 50°C.

- The subsequent reductive alkylation is preferably carried out in a suitable solvent such as methanol, methanol/water, methanol/water/ammonia, ethanol, ether, tetrahydrofuran,
30 dioxane or dimethylformamide, optionally with the addition of an acid such as hydrochloric acid in the presence of catalytically activated hydrogen, e.g. hydrogen in the presence of Raney nickel, platinum or palladium/charcoal, or in the presence of a metal

hydride such as sodium borohydride, sodium cyanoborohydride, lithium borohydride or lithium aluminium hydride at temperatures between 0 and 100°C, preferably at temperatures between 20 and 80°C.

5 The subsequent acylation is preferably carried out in a solvent such as methylene chloride, diethylether, tetrahydrofuran, toluene, dioxane, acetonitrile, dimethylsulphoxide or dimethylformamide, optionally in the presence of an inorganic or a tertiary organic base, preferably at temperatures between 20°C and the boiling temperature of the solvent used. The acylation with a corresponding acid is preferably carried out in the presence of
10 a dehydrating agent, e.g. in the presence of isobutyl chloroformate, tetraethyl orthocarbonate, trimethyl orthoacetate, 2,2-dimethoxypropane, tetramethoxysilane, thionylchloride, trimethylchlorosilane, phosphorus trichloride, phosphorus pentoxide, N,N'-dicyclohexylcarbodiimide, N,N'-dicyclohexyl-carbodiimide/N-hydroxysuccinimide, N,N'-dicyclohexylcarbodiimide/1-hydroxy-benzotriazole, 2-(1H-benzotriazol-1-yl)-
15 1,1,3,3-tetramethyluronium-tetrafluoroborate, 2-(1H-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium-tetrafluoroborate/1-hydroxy-benzotriazole, N,N'-carbonyldiimidazole or triphenylphosphine/carbon tetrachloride, and optionally with the addition of a base such as pyridine, 4-dimethylamino-pyridine, N-methyl-morpholine or triethylamine, conveniently at temperatures between 0 and 150°C,
20 preferably at temperatures between 0 and 100°C, and the acylation with a corresponding reactive compound such as an anhydride, ester, imidazolide or halide thereof is optionally carried out in the presence of a tertiary organic base such as triethylamine, N-ethyl-diisopropylamine or N-methyl-morpholine at temperatures between 0 and 150°C, preferably at temperatures between 50 and 100°C.

25

The subsequent esterification or amidation is expediently carried out by reacting a corresponding reactive carboxylic acid derivative with a corresponding alcohol or amine as described hereinbefore.

30 The subsequent reduction of a nitro group is preferably carried out by hydrogenolysis, e.g. with hydrogen in the presence of a catalyst such as palladium/charcoal or Raney nickel in a solvent such as methanol, ethanol, ethyl acetate, dimethylformamide,

dimethylformamide/acetone or glacial acetic acid, optionally with the addition of an acid such as hydrochloric acid or glacial acetic acid at temperatures of between 0 and 50°C, but preferably at ambient temperature, and at a hydrogen pressure of 1 to 7 bar, but preferably 3 to 5 bar.

5

The subsequent reduction of a cyano group is preferably carried out by hydrogenolysis, e.g. with hydrogen in the presence of a catalyst such as palladium/charcoal or Raney nickel in a solvent such as methanolic ammonia, ethanolic ammonia, ethyl acetate, dimethylformamide, dimethylformamide/acetone, dichloromethane or glacial acetic acid, optionally with the addition of an acid such as hydrochloric acid or glacial acetic acid at temperatures of between 0 and 50°C, but preferably at ambient temperature, and at a hydrogen pressure of 1 to 7 bar, but preferably 3 to 5 bar.

In the reactions described hereinbefore, any reactive groups present such as carboxy, amino, alkylamino or imino groups may be protected during the reaction by conventional protecting groups which are cleaved again after the reaction.

For example, a protecting group for a carboxyl group may be a trimethylsilyl, methyl, ethyl, tert.butyl, benzyl or tetrahydropyranyl group and

20

protecting groups for an amino, alkylamino or imino group may be an acetyl, trifluoroacetyl, benzoyl, ethoxycarbonyl, tert.butoxycarbonyl, benzyloxycarbonyl, benzyl, methoxybenzyl or 2,4-dimethoxybenzyl group and additionally, for the amino group, a phthalyl group.

25

Any protecting group used is optionally subsequently cleaved for example by hydrolysis in an aqueous solvent, e.g. in water, isopropanol/water, tetrahydrofuran/water or dioxane/water, in the presence of an acid such as trifluoroacetic acid, hydrochloric acid or sulphuric acid or in the presence of an alkali metal base such as lithium hydroxide, sodium hydroxide or potassium hydroxide, at temperatures between 0 and 100°C, preferably at temperatures between 10 and 50°C.

30

However, a benzyl, methoxybenzyl or benzyloxycarbonyl group is cleaved, for example, hydrogenolytically, e.g. with hydrogen in the presence of a catalyst such as palladium/charcoal in a solvent such as methanol, ethanol, ethyl acetate, dimethylformamide, dimethylformamide/acetone or glacial acetic acid, optionally with
5 the addition of an acid such as hydrochloric acid or glacial acetic acid at temperatures between 0 and 50°C, but preferably at ambient temperature, and at a hydrogen pressure of 1 to 7 bar, but preferably 3 to 5 bar.

A methoxybenzyl group may also be cleaved in the presence of an oxidising agent such
10 as cerium(IV)ammonium nitrate in a solvent such as methylene chloride, acetonitrile or acetonitrile/water at temperatures of between 0 and 50°C, but preferably at ambient temperature.

A 2,4-dimethoxybenzyl group, however, is preferably cleaved in trifluoroacetic acid in
15 the presence of anisole.

A tert.butyl or tert.butyloxycarbonyl group is preferably cleaved by treating with an acid such as trifluoroacetic acid or hydrochloric acid, optionally using a solvent such as methylene chloride, dioxane, ethyl acetate or ether.
20

A phthalyl group is preferably cleaved in the presence of hydrazine or a primary amine such as methylamine, ethylamine or n-butylamine in a solvent such as methanol, ethanol, isopropanol, toluene/water or dioxane at temperatures between 20 and 50°C.

Moreover, chiral compounds of general formula I obtained may be resolved into their
25 enantiomers and/or diastereomers.

Thus, for example, the compounds of general formula I obtained which occur as racemates may be separated by methods known per se (cf. Allinger N. L. and Eliel E. L.
30 in "Topics in Stereochemistry", Vol. 6, Wiley Interscience, 1971) into their optical antipodes and compounds of general formula I with at least 2 asymmetric carbon atoms may be resolved into their diastereomers on the basis of their physical-chemical

differences using methods known per se, e.g. by chromatography and/or fractional crystallisation, and, if these compounds are obtained in racemic form, they may subsequently be resolved into the enantiomers as mentioned above.

5 The enantiomers are preferably separated by column separation on chiral phases or by recrystallisation from an optically active solvent or by reacting with an optically active substance which forms salts or derivatives such as e.g. esters or amides with the racemic compound, particularly acids and the activated derivatives or alcohols thereof, and separating the diastereomeric mixture of salts or derivatives thus obtained, e.g. on the
10 basis of their differences in solubility, whilst the free antipodes may be released from the pure diastereomeric salts or derivatives by the action of suitable agents. Optically active acids in common use are e.g. the D- and L-forms of tartaric acid or dibenzoyltartaric acid, di-o-tolytartaric acid, malic acid, mandelic acid, camphorsulphonic acid, glutamic acid, N-acetylglutamic acid, aspartic acid, N-acetylaspatic acid or quinic acid. An optically
15 active alcohol may be for example (+)- or (-)-menthol and an optically active acyl group in amides, for example, may be a (+)- or (-)-menthyloxycarbonyl group.

Furthermore, the compounds of formula I obtained may be converted into the salts thereof, particularly for pharmaceutical use into the physiologically acceptable salts with
20 inorganic or organic acids. Acids which may be used for this purpose include for example hydrochloric acid, hydrobromic acid, sulphuric acid, phosphoric acid, fumaric acid, succinic acid, lactic acid, citric acid, tartaric acid, maleic acid or methanesulphonic acid.

Moreover, if the new compounds of formula I thus obtained contain a carboxy group,
25 they may subsequently, if desired, be converted into the salts thereof with inorganic or organic bases, particularly for pharmaceutical use into the physiologically acceptable salts thereof. Suitable bases for this purpose include for example sodium hydroxide, potassium hydroxide, cyclohexylamine, ethanolamine, diethanolamine and triethanolamine.

30

The compounds of general formulae II to V used as starting materials are known from the literature in some cases or may be obtained by methods known from the literature or are

described in the Examples. For example, the compounds of general formula IV are described in German Patent Application 198 24 922.5 of 4th June 1998.

5 As already mentioned hereinbefore, the new compounds of general formula I wherein R₁ denotes a hydrogen atom or a prodrug group have valuable pharmacological properties, particularly an inhibiting effect on the proliferation of cultivated human cells, especially tumour cells, but also on the proliferation of other cells, particularly endothelial cells, e.g. in angiogenesis.

10 For example, the compounds listed in Table 1 were tested for their biological properties as follows:

Test 1

15 Inhibition of the proliferation of cultivated human tumour cells

Cells of the Leiomyosarcoma tumour cell line SK-UT-1B or non-small-cell lung tumour cell line NCI-H460 (obtained from the American Type Culture Collection (ATCC)) were cultivated in Minimum Essential Medium with non-essential amino acids (Gibco), supplemented with sodium pyruvate (1 mMol), glutamine (2 mMol) and 10% foetal calf
20 serum (Gibco) or RPMI1640 Medium (Gibco) and 10% foetal calf serum (Gibco) and harvested in the logarithmic growth phase. Then the SK-UT-1B cells were placed in Cytostar® multi-well plates (Amersham) at a density of 4000 cells per well or 3000 cells per well for NCI-H460 cells and incubated overnight in an incubator. Various concentrations of the compounds (dissolved in DMSO; final concentration: 0.1%) were
25 added to the cells. After 48 hours' incubation, ¹⁴C-thymidine (Amersham) was added to each well and incubation was continued for a further 24 hours. The quantity of ¹⁴C-thymidine which was incorporated into the tumour cells in the presence of the inhibitor and which represents the number of cells in the S phase was measured in a Wallace 1450 Microbeta Liquid Scintillation Counter. IC₅₀ values for the inhibition of the
30 proliferation (= inhibition of incorporated ¹⁴C-thymidine) were calculated, correcting for the background radiation. All the measurements were done twice.

Test 2

In vivo effects on tumour-bearing nude mice

10⁶ cells [SK-UT-1B, or non-small cell lung tumour NCI-H460 (obtained from ATCC)]
 5 in a volume of 0.1 ml were injected subcutaneously into male and/or female nude mice
 (NMRI nu/nu; 25-35 g; N = 10-20); alternatively, small fragments of SK-UT-1B or NCI-
 H460 cell clumps were implanted subcutaneously. One to three weeks after injection or
 implantation an inhibitor was administered orally (by oesophageal tube) daily for a period
 of 2 to 4 weeks. The tumour size was measured three times a week using a digital sliding
 10 gauge. The effect of a compound on the tumour growth was determined as a percentage
 inhibition compared with a control group treated with placebo.

The following Table contains the results obtained with the *in vitro* Test 1 (++ denotes
 <0.01 μ M, + denotes 0.01-1.0 μ M):

15

Compound (Example No.)	Inhibition of SKUT-1B proliferation
2	+
4	++
9	+
12	+
20	+
22	+
23	+
31	++
36	++
42	+
56	++
58	+
66	++

70	+
71	+
72	+
80	++
88	+
98	+
99	++
101	++
104	++
112	++
117	+
120	++
134	++
142	+
143	+
144	+
145	+
158	+
164	+
186	++
207	+

In view of their biological properties, the new compounds of general formula I, their isomers and their physiologically acceptable salts are suitable for treating conditions characterised by excessive or anomalous cell proliferation.

5

Such diseases include (without any claim to completeness): viral infections (e.g. HIV and Kaposi's sarcoma); inflammation and autoimmune diseases (e.g. colitis, arthritis, Alzheimer's disease, glomerulonephritis and wound healing); bacterial, fungal and/or parasitic infections; leukaemias, lymphoma and solid tumours; skin diseases (e.g. psoriasis); bone diseases; cardiovascular diseases (e.g. restenosis and hypertrophy).

10

The new compounds may be used for the short-term or long-term treatment of the abovementioned conditions, possibly in conjunction with other state-of-the-art compounds such as other cytostatics.

5

The dosage required to achieve the desired effect is expediently from 0.1 to 30 mg/kg, preferably 0.3 to 10 mg/kg, by intravenous route and 0.1 to 100 mg/kg, preferably 0.3 to 30 mg/kg by oral route, in each case 1 to 4 times a day. For this purpose, the compounds of formula I prepared according to the invention, optionally combined with other active substances, may be formulated with one or more inert conventional carriers and/or diluents, e.g. with corn starch, lactose, glucose, microcrystalline cellulose, magnesium stearate, polyvinylpyrrolidone, citric acid, tartaric acid, water, water/ethanol, water/glycerol, water/sorbitol, water/polyethyleneglycol, propyleneglycol, cetylstearylalcohol, carboxymethylcellulose or fatty substances such as hard fat or suitable mixtures thereof, to produce conventional galenic preparations such as plain or coated tablets, capsules, powders, suspensions or suppositories or as solutions for injections or infusions.

10

15

The Examples which follow are intended to illustrate the invention without restricting it:

20

Abbreviations used:

CDI	N,N'-carbonyldiimidazole
DMF	dimethylformamide
25 DMSO	dimethylsulphoxide
TBTU	O-(benzotriazol-1-yl)-N,N,N'-N'-bis(tetramethylene)-uronium hexafluorophosphate
THF	tetrahydrofuran

30 Preparation of the starting compounds:

Example I

4-[N-Acetyl-N-(2-trifluoroacetyl-aminoethyl)-amino]-aniline

a. 4-(2-tert.Butoxycarbonylamino-ethylamino)-nitrobenzene

5 4.2 g (29.7 mmol) of N-tert.butoxycarbonyl-ethylenediamine, 5.0 g (31.2 mmol) of 4-fluoro-nitrobenzene and 7.0 g (50.6 mmol) of potassium carbonate are stirred in 25 ml of DMSO for 9 hours at 60°C. After cooling the mixture is diluted with water and extracted with ethyl acetate. The combined organic extracts are dried and evaporated down. The residue is stirred with petroleum ether, decanted off and evaporated down again. The
10 product is stirred with ether and suction filtered.

Yield: 3.2 g (38 % of theory),

Melting point: 119°C

R_f value: 0.5 (silica gel; toluene/ethyl acetate = 7:3)

C₁₃H₁₉N₃O₄ (281.31)

15 Mass spectrum: (M-H)⁻ = 280

b. 4-(2-trifluoroacetyl-amino-ethylamino)-nitrobenzene

1.5 g (5.3 mmol) of 4-(2-tert.butoxycarbonylamino-ethylamino)-nitrobenzene are stirred in 15 ml of trifluoroacetic acid for 3 hours at ambient temperature. Then 0.8 ml (5.7
20 mmol) of trifluoroacetic acid anhydride are added while cooling with ice. The reaction is left overnight to come up to ambient temperature. It is then evaporated down, diluted with water and made alkaline with sodium hydrogen carbonate. The crude product is suction filtered and purified by chromatography (silica gel; dichloromethane/methanol = 98:2).

25 Yield: 1.2 g (81 % of theory),

R_f value: 0.5 (silica gel; dichloromethane/methanol = 19:1)

C₁₀H₁₀F₃N₃O₃ (277.21)

Mass spectrum: (M-H)⁻ = 276

30 c. 4-[N-Acetyl-N-(2-trifluoroacetyl-amino-ethyl)-amino]-nitrobenzene

0.6 g (2.1 mmol) of 4-(2-trifluoroacetyl-amino-ethylamino)-nitrobenzene are dissolved in 10 ml of glacial acetic acid and after the addition of 2 ml (21.2 mmol) of acetic acid

anhydride stirred for 5 hours at 80°C and overnight at ambient temperature. The solvent is distilled off, the residue is made alkaline with sodium hydrogen carbonate and extracted with ethyl acetate. The combined organic extracts are dried and evaporated down.

5 Yield: 0.7 g (97 % of theory),

R_f value: 0.4 (silica gel; dichloromethane/methanol = 19:1)

C₁₂H₁₂F₃N₃O₄ (319.24)

Mass spectrum: (M-H)⁻ = 318

10 d. 4-[N-acetyl-N-(2-trifluoroacetyl-amino-ethyl)-amino]-aniline

0.7 g (2.1 mmol) of 4-[N-acetyl-N-(2-trifluoroacetyl-amino-ethyl)-amino]-nitrobenzene are dissolved in 20 ml of methanol and after the addition of 100 mg of 10% palladium on activated charcoal hydrogenated with hydrogen for 3 hours. Then the catalyst is filtered off and evaporated down.

15 Yield: 0.6 g (91 % of theory),

R_f value: 0.7 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₂H₁₄F₃N₃O₂ (289.26)

Mass spectrum: (M-H)⁻ = 288, (M+Na)⁺ = 312

20 The following compounds were prepared analogously to Example I :

(1) 4-[N-(2-dimethylamino-ethyl)-N-acetyl-amino]-aniline

R_f value: 0.3 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₂H₁₉N₃O (221.31)

25 Mass spectrum: (M+H)⁺ = 222

(2) 4-[N-(2-acetyl-amino-ethyl)-N-acetyl-amino]-aniline

R_f value: 0.4 (silica gel; ethyl acetate/methanol = 8:2)

C₁₂H₁₇N₃O₂ (235.28)

30 Mass spectrum: (M+Na)⁺ = 258, (M-H)⁻ = 234

(3) 4-[N-(2-acetyl-amino-ethyl)-N-propionyl-amino]-aniline

R_f value: 0.4 (silica gel; ethyl acetate/methanol = 9:1)

(4) [N-(2-propionylamino-ethyl)-N-propionyl-amino]-aniline

R_f value: 0.5 (silica gel; ethyl acetate/methanol = 9:1)

5

(5) 4-{N-[2-(N-acetyl-N-methyl-amino)-ethyl]-N-propionyl-amino}-aniline

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 19:1:0.1)

C₁₄H₂₁N₃O₂ (263.34)

Mass spectrum: (M+Na)⁺ = 286

10

(6) 4-{N-[2-(N-acetyl-N-methyl-amino)-ethyl]-N-acetyl-amino}-aniline

R_f value: 0.3 (silica gel; ethyl acetate/methanol = 9:1)

C₁₃H₁₉N₃O₂ (249.31)

Mass spectrum: (M-H)⁻ = 248, (M+Na)⁺ = 272

15

(7) 4-(dimethylaminocarbonylmethylamino)-aniline

R_f value: 0.6 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₀H₁₅N₃O (193.25)

Mass spectrum: (M+H)⁺ = 194, (M+Na)⁺ = 216

20

(8) 4-(N-ethoxycarbonylmethyl-N-acetyl-amino)-aniline

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₂H₁₆N₂O₃ (236.27)

Mass spectrum: (M-H)⁻ = 235, (M+Na)⁺ = 259

25

(9) 4-[N-(3-dimethylamino-propyl)-N-propionyl-amino]-aniline

R_f value: 0.2 (silica gel; dichloromethane/methanol/ammonia = 8.5:1.5:0.15)

C₁₄H₂₃N₃O (249.36)

Mass spectrum: (M-H)⁻ = 248, (M+H)⁺ = 250

30

Example II

4-[N-(2-benzyloxycarbonylamino-ethyl)-N-acetyl-amino]-aniline

450 mg (1.26 mmol) of 4-[N-(2-benzyloxycarbonylamino-ethyl)-N-acetyl-amino]-nitrobenzene (prepared analogously to Example I) are dissolved in 20 ml of methanol and after the addition of 100 mg of Lindlar catalyst hydrogenated for 2 hours with
5 hydrogen. The catalyst is filtered off, the solution is evaporated down.

Yield: 410 mg (99 % of theory),

R_f value: 0.4 (silica gel; ethyl acetate/dichloromethane = 7:3)

C₁₈H₂₁N₃O₃ (327.38)

Mass spectrum: (M+Na)⁺ = 350, (M-H)⁻ = 326

10

The following compounds were prepared analogously to Example II:

(1) 4-{N-[2-(N-benzyl-N-methyl-amino)-ethyl]-N-acetyl-amino}-aniline

R_f value: 0.7 (silica gel; ethyl acetate/methanol/ammonia = 9:1:0.1)

15 C₁₈H₂₃N₃O (297.40)

Mass spectrum: (M+H)⁺ = 298, (M-H)⁻ = 296

(2) 4-{N-[2-(N-benzyl-N-methyl-amino)-ethyl]-N-propionyl-amino}-aniline

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

20 C₁₉H₂₅N₃O (311.43)

Mass spectrum: (M+H)⁺ = 312

Example III

25 4-[N-(2-trifluoroacetyl-amino-ethyl)-N-methylsulphonyl-amino]-aniline

a. 4-(N-ethoxycarbonylmethyl-N-methylsulphonyl-amino)-nitrobenzene

20 g (92.5 mmol) of 4-(methylsulphonylamino)-nitroaniline are dissolved in 155 ml of DMSO and while cooling with ice 11.7 (104 mmol) of potassium tert. butoxide are
30 added. After 1 hour 13.5 ml (121 mmol) of ethyl bromoacetate are added. The mixture is stirred for 18 hours at ambient temperature and the reaction solution is then poured onto

ice water. It is extracted with ethyl acetate. The organic phase is washed with water, dried and freed from the solvent *in vacuo*. The residue is triturated with petroleum ether.

Yield: 27.1 g (97 % of theory),

Melting point: 73-75°C

5 R_f value: 0.8 (silica gel; dichloromethane/ethyl acetate = 9:1)

C₁₁H₁₄N₂O₆S (302.31)

Mass spectrum: (M+Na)⁺ = 325, (M-H)⁻ = 301

b. 4-(N-carboxymethyl-N-methylsulphonyl-amino)-nitrobenzene

10 26.8 g (88.6 mmol) of 4-(N-ethoxycarbonylmethyl-N-methylsulphonyl-amino)-nitrobenzene are suspended in 320 ml of ethanol and combined with 268 ml of 1 N sodium hydroxide solution. The mixture is stirred for one hour at ambient temperature and then 268 ml of 1 N hydrochloric acid are added. The precipitate formed is suction filtered, washed with a little ethanol and ether, and dried *in vacuo*.

15 Yield: 21.9 g (90% of theory),

Melting point: 215-218°C

R_f value: 0.6 (silica gel; dichloromethane/methanol/glacial acetic acid = 9:1:0.1)

C₉H₁₀N₂O₆S (274.25)

Mass spectrum: (M-H)⁻ = 273

20

c. 4-(N-aminocarbonylmethyl-N-methylsulphonyl-amino)-nitrobenzene

2.5 g (15.4 mmol) of CDI are added to a solution of 3 g (10.9 mmol) of 4-(N-carboxymethyl-N-methylsulphonyl-amino)-nitrobenzene in 30 ml of DMF. The mixture is stirred for one hour at ambient temperature. Then NH₃ is piped in at 0°C over a period
25 of 10 min. After 2 hours' stirring at ambient temperature 100 ml of water are added. The mixture is extracted with ethyl acetate, the organic phase is washed with water, dried over magnesium sulphate and evaporated to dryness. The residue is stirred with water, suction filtered and washed with ether.

Yield: 2.3 g (78% of theory),

30 Melting point: 160°C

R_f value: 0.5 (silica gel; ethyl acetate/dichloromethane = 3:2)

d. 4-[N-(2-aminoethyl)-N-methylsulphonyl-amino]-nitrobenzene

2.3 g (8.4 mmol) of 4-(N-aminocarbonylmethyl-N-methylsulphonyl-amino)-nitrobenzene are refluxed in 35 ml (35 mmol) of borane-THF (1 M solution in THF) 7 hours. Then 30 ml of 6 N hydrochloric acid are added, and the mixture is refluxed for another 8 hours.

5 The solvent is distilled off, the residue is mixed with water and extracted with ethyl acetate. The aqueous phase is made alkaline with potassium carbonate and extracted with dichloromethane. The organic phase is separated off, dried and evaporated down.

Yield: 1.7 g (77 % of theory),

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

10 C₉H₁₃N₃O₄S (259.29)

Mass spectrum: (M+H)⁺ = 260, (M-H)⁻ = 258

e. 4-[N-(2-trifluoroacetyl-amino-ethyl)-N-methylsulphonyl-amino]-aniline

Prepared analogously to Example Ib by reacting 4-[N-(2-aminoethyl)-N-methylsulphonyl-amino]-nitrobenzene with trifluoroacetic acid anhydride in
15 trifluoroacetic acid followed by catalytic reduction analogously to Example Id with 10% palladium/charcoal in methanol.

Yield: 76 % of theory,

R_f value: 0.6 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

20

The following compounds were prepared analogously to Example III:

(1) 4-(N-ethoxycarbonylmethyl-N-ethylsulphonyl-amino)-aniline

25 R_f value: 0.5 (silica gel; petroleum ether/ethyl acetate = 4:6)

Melting point: 78°C

C₁₂H₁₈N₂O₄S (286.35)

Mass spectrum: (M+Na)⁺ = 309, (2M+Na)⁺ = 593

30 (2) 4-{N-[2-(N-acetyl-N-methyl-amino)-ethyl]-N-methylsulphonyl-amino}-aniline

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 19:1:0.1)

(3) 4-[N-(2-acetylamino-ethyl)-N-methylsulphonyl-amino]-aniline

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 19:1:0.1)

C₁₁H₁₇N₃O₃S (271.34)

Mass spectrum: (M+H)⁺ = 272, (M+Na)⁺ = 294

5

(4) 4-{N-[2-(N-acetyl-N-methyl-amino)-ethyl]-N-ethylsulphonyl-amino}-aniline

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 19:1:0.1)

Melting point: 140°C

C₁₃H₂₁N₃O₃S (299.39)

10 Mass spectrum: M⁺ = 299

(5) 4-[N-(2-acetylamino-ethyl)-N-ethylsulphonyl-amino]-aniline

R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 19:1:0.1)

C₁₂H₁₉N₃O₃S (285.36)

15 Mass spectrum: (M-H)⁻ = 284, (M+Na)⁺ = 308

(6) 4-{N-[2-(N-methyl-N-trifluoroacetyl-amino)-ethyl]-N-methylsulphonyl-amino}-aniline

R_f value: 0.5 (silica gel; dichloromethane/ethyl acetate = 9:1)

20

Example IV

4-[N-(2-dimethylamino-ethyl)-N-phenylsulphonyl-amino]-aniline

25 a. N-(2-dimethylamino-ethyl)-phenylsulphonamide

2.8 g (30 mmol) of N,N-dimethylethylenediamine are placed in 100 ml of dichloromethane and 8.3 ml (60 mmol) of triethylamine. While cooling with ice a solution of 3.9 ml (30 mmol) of benzenesulphonic acid chloride in 100 ml of dichloromethane is added dropwise and the mixture is stirred overnight at ambient temperature. Water is added and the mixture is extracted with dichloromethane. The organic phase is dried and evaporated down.

30

Yield: 6.8 g (99 % of theory),

R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₀H₁₆N₂O₂S (228.23)

Mass spectrum: (M-H)⁻ = 227, (M+H)⁺ = 229

5 b. 4-[N-(2-dimethylamino-ethyl)-N-phenylsulphonyl-amino]-nitrobenzene

6.8 g (29.8 mmol) of N-(2-dimethylamino-ethyl)-phenylsulphonamide are dissolved in 100 ml of DMF and combined with 1.3 g (30 mmol) of sodium hydride (55% in oil). The mixture is stirred for one hour at ambient temperature. Then 4.2 g (29.8 mmol) of 4-fluoro-nitrobenzene are added, and stirring is continued for another 16 hours. After the addition of 300 ml of water the mixture is extracted with ethyl acetate. The organic phase is washed with water, dried and evaporated down. The residue is acidified with 1 N hydrochloric acid and washed with ethyl acetate. The aqueous phase is then made basic again with sodium hydroxide solution and extracted with ethyl acetate. The organic phase is dried and evaporated down.

15 Yield: 6.0 g (58 % of theory),

R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₆H₁₉N₃O₄S (349.41)

Mass spectrum: (M-H)⁻ = 348, (M+H)⁺ = 350

20 c. 4-[N-(2-dimethylaminoethyl)-N-phenylsulphonyl-amino]-aniline

Prepared analogously to Example Id by catalytic hydrogenation of 6 g (17.2 mmol) of 4-[N-(2-dimethylamino-ethyl)-N-phenylsulphonyl-amino]-nitrobenzene.

Yield: 5.5 g (99 % of theory),

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

25 C₁₆H₂₁N₃O₂S (319.43)

Mass spectrum: (M+H)⁺ = 320

The following compounds were prepared analogously to Example IV:

30 (1) 4-[N-(2-dimethylamino-ethyl)-N-propylsulphonyl-amino]-aniline

R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₃H₂₃N₃O₂S (285.41)

Mass spectrum: $(M+H)^+ = 286$, $(M-H)^- = 284$

(2) 4-[N-(2-dimethylamino-ethyl)-N-butylsulphonyl-amino]-aniline

R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

5 C₁₄H₂₅N₃O₂S (299.43)

Mass spectrum: $(M+H)^+ = 300$

(3) 4-[N-(3-dimethylamino-propyl)-N-methylsulphonyl-amino]-aniline

Melting point: 112-113°C

10 R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₂H₂₁N₃O₂S (271.38)

Mass spectrum: $(M+H)^+ = 272$, $(M+Na)^+ = 294$

(4) 4-[N-(2-dimethylamino-ethyl)-N-benzylsulphonyl-amino]-aniline

15 R_f value: 0.3 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₇H₂₃N₃O₂S (333.46)

Mass spectrum: $(M+H)^+ = 334$, $(M+Na)^+ = 356$

(5) 3-chloro-4-[N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-aniline

20 Melting point: 145-148°C

R_f value: 0.5 (silica gel; dichloromethane/ethanol/ammonia = 5:1:0.01)

C₁₁H₁₈ClN₃O₂S (291.80)

Mass spectrum: $(M+H)^+ = 294, 292$, $(M-H)^- = 292, 290$

25 (6) 3-amino-4-[N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-aniline

R_f value: 0.3 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₁H₂₀N₄O₂S (272.37)

Mass spectrum: $(M+H)^+ = 273$

30 (7) 4-[N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-aniline

R_f value: 0.3 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

Melting point: 147-148°C

$C_{11}H_{19}N_3O_2S$ (257.36)

Mass spectrum: $(M+H)^+ = 258$, $(M+Na)^+ = 280$

Example V

5

3-[N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-aniline

a. 3-[N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-nitrobenzene

5 g (23.1 mmol) of 3-methylsulphonylamino-nitrobenzene are dissolved in 50 ml of
10 DMSO and combined with 6.5 g (58 mmol) of potassium tert. butoxide while cooling
with ice. The solution thus obtained is added dropwise to a solution of 5 g (34.7 mmol) of
2-chloro-N,N-dimethyl-ethylamine in 30 ml of DMSO. The mixture is stirred for 2 hours
at ambient temperature and then heated for 6 hours to 100 °C. After cooling to ambient
temperature 400 ml of water are added. The mixture is extracted with ethyl acetate. Water
15 and 1 N hydrochloric acid are added to the combined organic phases until an acid
reaction is obtained. The aqueous phase is washed with ethyl acetate. Then the aqueous
phase is made alkaline with sodium carbonate and the product is extracted with ethyl
acetate. Drying the combined organic phases over magnesium sulphate and eliminating
the solvents *in vacuo* yields the product as a red oil.

20 Yield: 2.07 g (31 % of theory),

R_f value: 0.3 (silica gel; ethyl acetate/methanol = 4:1)

$C_{11}H_{17}N_3O_4S$ (287.34)

Mass spectrum: $(M-H)^- = 286$, $(M+H)^+ = 288$

25 b. 3-[N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-aniline

Prepared analogously to Example 1d by catalytic hydrogenation of 1.9 g (6.8 mmol) of 3-
[N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-nitrobenzene over
palladium/charcoal.

Yield: 1.8 g (99% of theory),

30 R_f value: 0.3 (silica gel; ethyl acetate/methanol/ NH_4OH = 8:2:0.1)

$C_{11}H_{19}N_3O_2S$ (257.36)

Mass spectrum: $(M-H)^- = 256$, $(M+H)^+ = 258$

Example VI

4-(4-benzyl-piperazinomethyl)-aniline

5

a. 4-(4-tert.butoxycarbonyl-piperazinomethyl)-nitrobenzene

A mixture of 10.6 g (57 mmol) of N-tert.butoxycarbonyl-piperazine, 10.8 g (62.7 mmol) of 4-nitrobenzylchloride, 23.8 ml (171 mmol) of triethylamine in 100 ml of dichloromethane is stirred for 12 hours at 70°C. After diluting with water the organic
10 phase is separated off, dried and evaporated down.

Yield: 19 g (99 % of theory),

Melting point: 83-84°C

R_f value: 0.7 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₆H₂₃N₃O₄ (321.38)

15 Mass spectrum: (M+H)⁺ = 322, (M-H)⁻ = 320

b. 4-piperazinomethyl-nitrobenzene-dihydrochloride

6.4 g (20 mmol) of 4-(4-tert.butoxycarbonyl-piperazinomethyl)-nitrobenzene are dissolved in 20 ml of dichloromethane and combined with 40 ml of ethyl acetate/HCl.

20 The reaction solution is diluted with ether, the precipitate formed is suction filtered as a crude product and then reacted further.

Yield: 5.4 g (92 % of theory),

Melting point: 257-258°C

R_f value: 0.3 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

25

c. 4-(4-benzylpiperazinomethyl)-nitrobenzene

The free base is produced from 1.5 g (5 mmol) of 4-piperazinomethyl-nitrobenzene-dihydrochloride by dissolving in 25 ml of 1 N sodium hydroxide solution, extracting with ethyl acetate and then eliminating the solvent *in vacuo*. The solid thus obtained is
30 combined with 2.5 ml of 2 N acetic acid, 0.5 ml (5.5 mmol) of benzaldehyde and 50 ml of methanol and, after the addition of 0.7 g (5 mmol) of sodium cyanoborohydride, stirred for 2 hours. Then the pH is adjusted to acid with 1 N hydrochloric acid and the reaction

solution is washed with ether. The aqueous phase is then made basic with sodium hydroxide solution. The product is extracted with ether, the combined ether extracts are dried and the solvent is eliminated *in vacuo*.

Yield: 1.3 g (84 % of theory),

5 R_f value: 0.6 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

$C_{18}H_{21}N_3O_2$ (311.39)

Mass spectrum: $(M+H)^+ = 312$

d. 4-(4-benzylpiperazinomethyl)-aniline

10 Prepared analogously to Example Id by catalytic hydrogenation of 1.3 g (4.2 mmol) of 4-(4-benzylpiperazinomethyl)-nitrobenzene over palladium/charcoal.

Yield: 1.2 g (87 % of theory),

Melting point: 88-89°C

$C_{18}H_{23}N_3$ (281.4)

15 Mass spectrum: $(M+H)^+ = 282$

Example VII

4-(4-tert.butoxycarbonyl-piperazinomethyl)-aniline

20

a. 4-(4-tert.butoxycarbonyl-piperazinomethyl)-nitrobenzene

10.6 g (57 mmol) of N-tert.butoxycarbonyl-piperazine are dissolved in 100 ml of dichloromethane and combined with 10.7 g (63 mmol) of 4-nitrobenzylchloride and 24 ml (171 mmol) of triethylamine. The mixture is refluxed for 12 hours. After cooling to ambient temperature the reaction solution is washed several times with water. The organic phase is dried over magnesium sulphate and then evaporated to dryness.

25

Yield: 17 g (99%) of theory

Melting point: 83-84°C

R_f value: 0.7 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

30 $C_{16}H_{23}N_3O_4$ (321.38)

Mass spectrum: $(M+H)^+ = 322$, $(M-H)^- = 320$

b. 4-(4-tert.butoxycarbonyl-piperazinomethyl)-aniline

Prepared analogously to Example Id by catalytic hydrogenation of 4-(4-tert.butoxycarbonyl-piperazinomethyl)-nitrobenzene with Raney nickel in ethyl acetate/methanol (1:1).

5 Melting point: 106-107°C

R_f value: 0.6 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₆H₂₅N₃O₂ (291.39)

Mass spectrum: (M+H)⁺ = 292, (M+Na)⁺ = 314

10 The following compounds were prepared analogously to Example VII:

(1) 4-(pyrrolidin-1-yl-methyl)-aniline

R_f value: 0.2 (silica gel; dichloromethane/methanol/NH₄OH = 5:1:0.01)

Melting point: 48-50°C

15

(2) 4-(4-methylpiperazinomethyl)-aniline

Melting point: 94-95°C

R_f value: 0.2 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

C₁₂H₁₉N₃ (205.31)

20 Mass spectrum: (M+H)⁺ = 206

(3) 3-(dimethylaminomethyl)-aniline

R_f value: 0.7 (silica gel; ethyl acetate)

Melting point: 43-46°C

25

(4) 4-(dimethylaminomethyl)-aniline

R_f value: 0.13 (silica gel; ethyl acetate/ethanol = 8:2)

(5) 4-(2-dimethylamino-ethyl)-aniline

30 R_f value: 0.3 (silica gel; dichloromethane/methanol/ammonia = 19:1:0.1)

Melting point: 40°C

C₁₀H₁₆N₂ (164.25)

Mass spectrum: $(M+H)^+ = 165$

(6) 4-(N-benzyl-N-methyl-aminomethyl)-aniline

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 10:1:0.01)

5 Melting point: 48-50°C

C₁₅H₁₈N₂ (226.32)

Mass spectrum: $(M+H)^+ = 227$

(7) 4-piperidinomethyl-aniline

10 R_f value: 0.2 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

Melting point: 88-89°C

(8) 4-(2,6-dimethylpiperidino-methyl)-aniline

R_f value: 0.3 (silica gel; dichloromethane/methanol/ammonia = 5:1:0.01)

15 Melting point: 112-115°C

(9) 4-(N-ethyl-N-methyl-aminomethyl)-aniline

R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 10:1:0.1)

C₁₀H₁₆N₂ (164.25)

20 Mass spectrum: $(M+H)^+ = 165$

(10) 4-[4-(3-trifluoromethylcarbonylamino-propyl)-piperidinomethyl]-aniline

R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 10:1:0.1)

C₁₇H₂₄F₃N₃O (343.40)

25 Mass spectrum: $(M+H)^+ = 344$

(11) 4-(N-tert.butoxycarbonyl-N-propyl-aminomethyl)-aniline

C₁₅H₂₄N₂O₂ (264.37)

Mass spectrum: $(M+Na)^+ = 287$

30

(12) 4-(N-tert.butoxycarbonyl-N-butyl-aminomethyl)-aniline

R_f value: 0.19 (silica gel; dichloromethane/methanol = 50:1)

$C_{16}H_{26}N_2O_2$ (278.40)

Mass spectrum: $(M+Na)^+ = 301$

(13) 4-(N-tert.butoxycarbonyl-N-ethyl-aminomethyl)-aniline

5 Melting point: 85°C

R_f value: 0.3 (silica gel; dichloromethane/methanol/ = 50:1)

$C_{14}H_{22}N_2O_2$ (250.34)

Mass spectrum: $(M+Na)^+ = 273$

10 Example VIII

4-(2-oxopiperidinomethyl)-aniline

6.4 g (42 mmol) of 4-nitrobenzaldehyde are dissolved in 150 ml of methanol and
15 combined with 4.9 g (42 mmol) of 5-aminovaleric acid and 1.8 g (29 mmol) of sodium
cyanoborohydride. The mixture is stirred for 18 hours at ambient temperature and then
carefully mixed with 20 ml of conc. hydrochloric acid. The solvent is eliminated *in*
vacuo, the residue is taken up in water and extracted with dichloromethane. The residue
obtained after evaporation is chromatographed on silica gel (dichloromethane/methanol,
20 4:1). A mixture of methyl 5-(4-nitrobenzylamino)-pentanoate and 4-(2-
oxopiperidinomethyl)-nitrobenzene is obtained which is dissolved in 100 methanol and
combined with 50 ml of 1 N sodium hydroxide solution. The mixture is stirred for one
hour at ambient temperature, 50 ml of 1 N hydrochloric acid are added and the reaction
solution is evaporated down to 100 ml. The aqueous phase thus obtained is extracted with
25 dichloromethane. The combined organic phases are dried over sodium sulphate and
evaporated to dryness.

The residue is hydrogenated analogously to Example Id over Raney nickel in methanol
under a hydrogen atmosphere of 3 bar for 11 hours.

Total yield: 2.2 g (26 % of theory),

30 R_f value: 0.63 (silica gel; dichloromethane/methanol = 9:1)

Example IX

4-(N-piperidinomethylcarbonyl-N-methyl-amino)-aniline

a. 4-(N-bromomethylcarbonyl-N-methyl-amino)-nitrobenzene

5 23.5 g (0.15 mol) of N-methyl-4-nitroaniline are dissolved in 400 ml of dioxane and combined with 22.2 g (0.3 mol) of lithium carbonate. Then 32.2 g (0.18 mol) of bromoacetyl bromide are added dropwise in such a way that the internal temperature does not exceed 33°C. After 18 hours' stirring the reaction solution is evaporated down to 100 ml, combined with 500 ml of water and stirred for 1 hour. The precipitate formed is
10 suction filtered, washed with water and dried. The crude product is stirred in 400 ml of ethyl acetate at 40°C. Then the insoluble matter is filtered off, the solution is evaporated down and the solid residue is triturated with ether.

Yield: 35 g (83 % of theory),

Melting point: 85-87°C

15

b. 4-(N-piperidinomethylcarbonyl-N-methyl-amino)-nitrobenzene

5.4 g (20 mmol) of 4-(N-bromomethylcarbonyl-N-methyl-amino)-nitrobenzene are dissolved in 100 ml of acetone and combined with 5.5 g (40 mmol) of potassium carbonate. 3 ml (30 mmol) of piperidine are slowly added dropwise and the mixture is
20 stirred for 18 hours at ambient temperature. The reaction solution is filtered, and the filtrate is evaporated to dryness. The residue is dissolved in ethyl acetate, washed with water, dried over magnesium sulphate and evaporated to dryness.

Yield: 5.6 g (99 % of theory).

25 c. 4-(N-piperidinomethylcarbonyl-N-methyl-amino)-aniline

Prepared analogously to Example Id by catalytic hydrogenation of 4-(N-piperidinomethylcarbonyl-N-methyl-amino)-nitrobenzene in methanol over palladium/charcoal.

Yield 4.95 g (99% of theory)

30

Example X

4-(tert.butoxycarbonylaminomethyl)-aniline

20 g (164 mmol) of 4-aminobenzylamine and 20.2 g (210 mmol) of triethylamine are dissolved in 100 ml of dioxane and 50 ml of water. 35.8 g (165 mmol) of di-tert.butyl-dicarbonate dissolved in 60 ml of dioxane are added to this solution while cooling with ice and the resulting mixture is stirred for 18 hours at ambient temperature. Then the solvent is distilled off *in vacuo*, the residue is distributed in ethyl acetate/water. The combined organic extracts are freed from solvent *in vacuo*. The crude product is heated in 200 ml of petroleum ether, cooled slowly with vigorous stirring and the crystalline product is removed by suction filtering.

Yield: 34.8 g (96 % of theory),
Melting point: 77-78°C

Example XI

4-(1H-imidazol-2-yl)-aniline

7.2 g (50 mmol) of 2-phenylimidazol are dissolved in 100 ml of conc. sulphuric acid. While cooling with ice 5.0 g (62 mmol) of ammonium nitrate are added in batches and the mixture is stirred for 2.5 hours. The reaction solution is then poured onto ice, made basic with conc. ammonia and the crystalline product is suction filtered. The nitro compound thus obtained is catalytically hydrogenated analogously to Example Id in DMF over palladium/charcoal.

Yield: 24 % of theory,
R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

Example XII

Pyridine-2-sulphonic acid chloride

5.0 g (45 mmol) of pyridine-2-thiol are dissolved in 40 ml of conc. hydrochloric acid. While the solution is cooled with ice, chlorine gas is piped in over a period of 2.5 hours. In order to destroy any excess gas a washing bottle containing 1 N sodium thiosulphate solution is attached. Then the reaction solution is poured onto ice water and extracted

with ether and dichloromethane. The organic phases are combined, dried and freed from solvent *in vacuo*. The crude product is further reacted immediately.

Yield: 8 g (100 % of theory).

5 Example XIII

Pyridine-3-sulphonic acid chloride hydrochloride

1 g (6.3 mmol) of pyridine-3-sulphonic acid and 1.4 g (6.7 mmol) of phosphorus pentachloride are stirred for 2 hours at 150°C. After cooling, excess phosphorus
10 pentachloride is eliminated *in vacuo*. The crude product is further reacted immediately.
Yield: 1.2 g (91 % of theory).

Preparation of the end products:

15 Example 1

(Z)-3-{1-[4-(N-acetyl-N-(2-aminoethyl)-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone

20 a. 1-acetyl-2-indolinone

13.3 g (0.1 mol) of 2-indolinone and 30 ml of acetic anhydride are stirred for 3 hours at 170°C. After cooling the mixture is combined with 150 ml of ice water, the crystalline product is suction filtered, washed with water and dried.

Yield: 16.6 g (95 % of theory),

25 Melting point: 129-130°C.

b. 1-acetyl-5-nitro-2-indolinone

0.5 g (2.8 mmol) of 1-acetyl-2-indolinone are placed in 4 ml of conc. sulphuric acid. At a temperature of -10 to -5°C, 0.3 g (3.4 mmol) of ammonium nitrate are added in batches.

30 After 45 minutes the mixture is poured onto ammonia/ice water, the crystalline precipitate is suction filtered, washed with water and dried. The crude product is recrystallised from 70 ml of cyclohexane.

Yield: 0.2 g (32 % of theory),

Melting point: 150-157°C

R_f value: 0.7 (silica gel; cyclohexane/ethyl acetate = 4:6)

5 c. 1-acetyl-5-amino-2-indolinone

30.0 g (136 mmol) of 1-acetyl-5-nitro-2-indolinone are dissolved in a mixture of 650 ml of dichloromethane and 650 ml of methanol and after the addition of 5 g of 10% palladium on activated charcoal the mixture is hydrogenated for 45 minutes with hydrogen. Then the catalyst is filtered off and evaporated down.

10 Yield: 22.4 g (87 % of theory),

Melting point: 177°C

R_f value: 0.7 (silica gel; ethyl acetate)

C₁₀H₁₀N₂O₂ (190.20)

Mass spectrum: (M-H)⁻ = 189, (M+Na)⁺ = 213

15

d. 1-acetyl-5-phenylsulphonylamino-2-indolinone

20.0 g (105 mmol) of 1-acetyl-5-amino-2-indolinone are placed in 200 ml of pyridine, combined with 15.3 ml (120 mmol) of benzenesulphonic acid chloride while cooling with ice and stirred for 2 hours. Then the mixture is poured onto 1.8 l of water and suction

20 filtered. The crude product is stirred into acetone, suction filtered and dried.

Yield: 30.5 g (88 % of theory),

Melting point: 245°C

R_f value: 0.5 (silica gel; dichloromethane/ethyl acetate = 9:1)

C₁₆H₁₄N₂O₄S (330.37)

25 Mass spectrum: (M-H)⁻ = 329, (M+Na)⁺ = 353

e. 1-acetyl-3-(1-ethoxy-1-phenyl-methylidene)-5-(N-acetyl-N-phenylsulphonyl-amino)-2-indolinone

8.0 g (24.2 mmol) of 1-acetyl-5-phenylsulphonylamino-2-indolinone are dissolved in 150 ml of acetic anhydride and after the addition of 20 ml (88.1 mmol) of triethyl orthobenzoate refluxed for 6 hours. The solvent is distilled off, the residue is triturated with ether, suction filtered and dried.

30

Yield: 7.8 g (64 % of theory),

Melting point: 237°C

R_f value: 0.7 (silica gel; dichloromethane/ethyl acetate = 19:1)

C₂₇H₂₄N₂O₆S (504.57)

5 Mass spectrum: M⁺ = 504

f. (Z)-3-{1-[4-(N-acetyl-N-(2-aminoethyl)-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone

A mixture of 0.5 g (1 mmol) of 1-acetyl-3-(1-ethoxy-1-phenyl-methylidene)-5-(N-acetyl-N-phenylsulphonyl-amino)-2-indolinone and 0.3 g (1.2 mmol) of 4-[N-acetyl-N-(2-trifluoroacetyl-amino-ethyl)-amino]-aniline are stirred in 5 ml of DMF for 6 hours at 120°C. After cooling to ambient temperature 5 ml of methanol and 3 ml (6 mmol) of 2 N sodium hydroxide solution are added, and the mixture is stirred for 30 minutes. The reaction mixture is diluted with 50 ml of water and the crystalline precipitate is suction
10 filtered and dried. The residue is chromatographed on silica gel
15 (dichloromethane/methanol/ammonia = 9:1:0.1).

Yield: 0.3 g (49 % of theory),

Melting point: 216°C

R_f value: 0.3 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

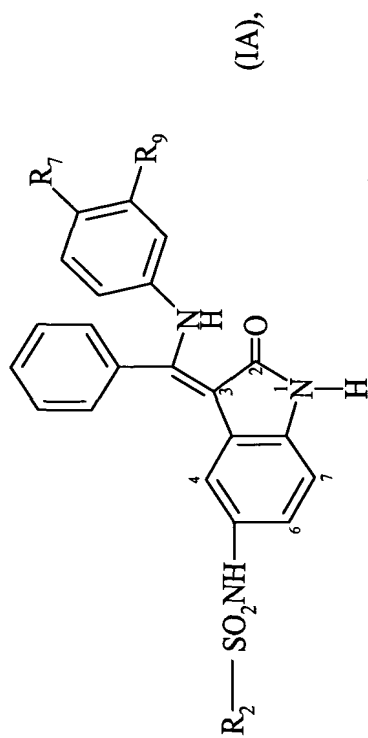
20 C₃₁H₂₉N₅O₄S (567.67)

Mass spectrum: (M-H)⁻ = 566, (M+H)⁺ = 568

Examples 2 to 97

25 Using the intermediate products prepared in Examples I to XIII, the compounds of formula IA of Examples 2 to 97 listed in Table I are prepared analogously to Example 1.

Table I



Example	R ₂	R ₇	R ₉	chemical name	Melting point (°C)
	phenyl	N-(2-aminoethyl)-N-methylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(2-aminoethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	245
	phenyl	N-(2-methylamino-ethyl)-N-methylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(2-methylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	227-229

phenyl	N-(2-dimethylamino-ethyl)-N-phenylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(2-dimethylaminoethyl)-N-phenylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	168-169
phenyl	N-(2-dimethylamino-ethyl)-N-propylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-propylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	137-138
phenyl	N-(2-dimethylamino-ethyl)-N-butylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-butylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	197-198
phenyl	4-benzyl-piperazino-methyl	H	(Z)-3-{1-[4-(4-benzyl-piperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	130 (decomp.)
phenyl	N-acetyl-N-(2-benzyl-oxycarbonylamino-ethyl)-amino	H	(Z)-3-{1-[4-(N-acetyl-N-(2-benzyl-oxycarbonylamino-ethyl)-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonyl-amino-2-indolinone	180
phenyl	4-methylpiperazino-methyl	H	(Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-	243-244

				phenylsulphonylamino-2-indolinone		
phenyl	morpholinomethyl	H		(Z)-3-{1-[4-(morpholinomethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	243-244	
phenyl	2-oxopiperidinomethyl	H		(Z)-3-{1-[4-(2-oxopiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	311-312	
phenyl	pyrrolidin-1-ylmethyl	H		(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	228-229	
phenyl	4-tert.butoxycarbonyl-piperazinomethyl	H		(Z)-3-{1-[4-(4-tert.butoxycarbonyl-piperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	160-161	
phenyl	N-methyl-N-formyl-amino	H		(Z)-3-{1-[4-(N-methyl-N-formyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	315-317	
phenyl	tert.butoxycarbonyl-amino	H		(Z)-3-[1-(4-tert.butoxycarbonylamino-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	96-98	

	phenyl	N-methyl-N-propionyl-amino	H	(Z)-3-{1-[4-(N-methyl-N-propionyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	208-210
	phenyl	acetylamino	H	(Z)-3-[1-(4-acetylamino-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	245-247
	phenyl	N-methyl-N-ethylsulphonyl-amino	H	(Z)-3-{1-[4-(N-methyl-N-ethylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	278-280
	phenyl	propionylamino	H	(Z)-3-[1-(4-propionylamino-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	254-256
	phenyl	N-methyl-N-acetyl-amino	H	(Z)-3-{1-[4-(N-methyl-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	283-285
	phenyl	N-acetyl-N-[2-(N-benzyl-N-methyl-amino)-ethyl]-amino	H	(Z)-3-{1-[4-(N-acetyl-N-(2-(N-benzyl-N-methyl-amino)-ethyl)-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	237
	phenyl	H	H	(Z)-3-(1-phenylamino-1-phenyl-methylidene)-5-phenylsulphonylamino-2-indolinone	283
	phenyl	chloro	H	(Z)-3-[1-(4-chlorophenylamino)-1-phenyl-	295

				methylidene]-5-phenylsulphonylamino-2-indolinone	
phenyl	N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	234	
phenyl	N-(2-dimethylamino-ethyl)-N-acetyl-amino	H	(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	202	
phenyl	N-piperidinomethyl-carbonyl-N-methyl-amino	H	(Z)-3-{1-[4-(N-piperidinomethylcarbonyl-N-methyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	269	
phenyl	H	N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino	(Z)-3-{1-[3-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	140	
phenyl	H	dimethylamino-methyl	(Z)-3-{1-[3-(N-(2-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene)-5-phenylsulphonylamino-2-indolinone	140	
phenyl	N-(2-acetyl-amino-ethyl)-N-acetyl-amino	H	(Z)-3-{1-[4-(N-(2-acetyl-amino-ethyl)-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	229	

				phenylsulphonylamino-2-indolinone		
phenyl	N-(2-acetylamino-ethyl)-N-propionyl-amino	H		(Z)-3-[1-[4-(N-(2-acetylamino-ethyl)-N-propionyl-amino)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	278	
phenyl	N-(2-propionylamino-ethyl)-N-propionyl-amino	H		(Z)-3-[1-[4-(N-(2-propionylamino-ethyl)-N-propionyl-amino)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	280	
phenyl	N-[2-(N-acetyl-N-methyl-amino)-ethyl]-N-methylsulphonyl-amino	H		(Z)-3-[1-[4-(N-(2-(N-acetyl-N-methyl-amino)-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	180 (decomp.)	
phenyl	N-(2-acetylamino-ethyl)-N-methylsulphonyl-amino	H		(Z)-3-[1-[4-(N-(2-acetylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	171	
phenyl	4-[N-[2-(N-acetyl-N-methyl-amino)-ethyl]-N-ethylsulphonyl-amino]	H		(Z)-3-[1-[4-(N-(2-(N-acetyl-N-methyl-amino)-ethyl)-N-ethylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	216	
phenyl	cyano	H		(Z)-3-[1-(4-cyanophenylamino)-1-phenyl-	291-293	

					methylidene]-5-phenylsulphonylamino-2-indolinone	
phenyl	dimethylaminomethyl	H			(Z)-3-[1-(4-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indole	255-256
phenyl	2-dimethylamino-ethyl	H			(Z)-3-[1-(4-(2-dimethylamino-ethyl)-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	302-303
phenyl	N-(2-acetyl-amino-ethyl)-N-ethylsulphonyl-amino	H			(Z)-3-{1-[4-(N-(2-acetyl-amino-ethyl)-N-ethylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	158
phenyl	acetylaminomethyl	H			(Z)-3-[1-(4-acetylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	289-290
phenyl	N-[2-(N-acetyl-l-N-methyl-amino)-ethyl]-N-acetyl-amino	H			(Z)-3-{1-[4-(N-(2-(N-acetyl-N-methyl-amino)-ethyl)-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	297
phenyl	methylsulphonylamin	H			(Z)-3-[1-(4-methylsulphonylamino-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	258-260

phenyl	N-methyl-N-methylsulphonylamino	H	(Z)-3-[1-(4-(N-methyl-N-methylsulphonylamino)-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	306-308
phenyl	ethylsulphonylamino	H	(Z)-3-[1-(4-ethylsulphonylamino-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	177-179
phenyl	N-[2-(N-acetyl-N-methyl-amino)-ethyl]-N-propionyl-amino	H	(Z)-3-[1-[4-(N-(2-(N-acetyl-N-methyl-amino)-ethyl)-N-propionyl-amino)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	250
phenyl	N-[2-(N-benzyl-N-methyl-amino)-ethyl]-N-propionyl-amino	H	(Z)-3-[1-[4-(N-(2-(N-benzyl-N-methyl-amino)-ethyl)-N-propionyl-amino)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	220
phenyl	dimethylamino-carbonylmethylamino	H	(Z)-3-[1-[4-(dimethylaminocarbonylmethylamino)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	230-231
phenyl	formylamino	H	(Z)-3-[1-(4-formylamino-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	305-307
phenyl	(2,6-dimethylpiperidino)-methyl	H	(Z)-3-[1-[4-((2,6-dimethylpiperidino)-methyl)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	144-145

				phenylsulphonylamino-2-indolinone		
phenyl	N-(dimethyl-aminomethylcarbonyl)-N-methyl-amino	H	(Z)-3-{1-[4-(N-dimethylaminomethylcarbonyl)-N-methyl-amino]-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	242		
phenyl	N-(2-dimethylamino-ethyl)-N-benzylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-benzylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	80 (decomp.)		
phenyl	2-propionylamino-ethylamino	H	(Z)-3-{1-[4-(2-propionylamino-ethylamino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	216		
phenyl	N-tert.butoxycarbonyl-N-propyl-aminomethyl	H	(Z)-3-{1-[4-(N-tert.butoxycarbonyl-N-propyl-aminomethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	215		
phenyl	N-tert.butoxycarbonyl-N-butyl-aminomethyl	H	(Z)-3-{1-[4-(N-tert.butoxycarbonyl-N-butyl-aminomethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	207		
phenyl	methyl	H	(Z)-3-[1-(4-methylphenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	192		

phenyl	N-methyl-N-ethyl-aminomethyl	H	(Z)-3-[1-(4-(N-methyl-N-ethyl-aminomethyl)-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	256
phenyl	N-methyl-N-piperidinomethylcarbonylamino	H	(Z)-3-[1-(4-(N-methyl-N-piperidinomethylcarbonylamino)-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	274-276
benzyl	dimethylaminomethyl	H	(Z)-3-{1-[4-dimethylaminomethyl-phenylamino]-1-phenyl-methylidene}-5-benzylsulphonylamino-2-indolinone	242-243
benzyl	pyrrolidin-1-ylmethyl	H	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl-methylidene}-5-benzylsulphonylamino-2-indolinone	228
benzyl	tert.butoxycarbonylaminomethyl	H	(Z)-3-[1-(4-tert.butoxycarbonylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-benzylsulphonylamino-2-indolinone	205
benzyl	(2,6-dimethylpiperidino)-methyl	H	(Z)-3-{1-[4-(2,6-dimethylpiperidino)-methyl]-phenylamino]-1-phenyl-methylidene}-5-benzylsulphonylamino-2-indolinone	140 (decomp.)

3-methoxy-phenyl	(2,6-dimethylpiperidino)-methyl	H	(Z)-3-{1-[4-((2,6-dimethylpiperidino)-methyl)-phenylamino]-1-phenyl-methylidene}-5-(3-methoxyphenylsulphonylamino)-2-indolinone	186
3-methoxy-phenyl	pyrrolidin-1-ylmethyl	H	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-methoxyphenylsulphonylamino)-2-indolinone	233
3-methoxy-phenyl	tert.butoxycarbonyl-aminomethyl	H	(Z)-3-[1-(4-tert.butoxycarbonylaminomethyl)-phenylamino]-1-phenyl-methylidene]-5-(3-methoxyphenylsulphonylamino)-2-indolinone	189
3-nitro-phenyl	pyrrolidin-1-ylmethyl	H	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-nitrophenylsulphonylamino)-2-indolinone	181
3-nitro-phenyl	tert.butoxycarbonyl-aminomethyl	H	(Z)-3-[1-(4-tert.butoxycarbonylaminomethyl)-phenylamino]-1-phenyl-methylidene]-5-(3-nitrophenylsulphonylamino)-2-indolinone	238°C (decomp.)
3-nitro-phenyl	(2,6-dimethylpiperidino)-methyl	H	(Z)-3-{1-[4-((2,6-dimethylpiperidino)-methyl)-phenylamino]-1-phenyl-methylidene}-5-(3-nitrophenylsulphonylamino)-2-indolinone	215

	2-cyano-phenyl	4-methylpiperazino-methyl	H	(Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(2-cyanophenylsulphonylamino)-2-indolinone	255 (decomp.)
	3-amino-carbonyl-phenyl	4-methylpiperazino-methyl	H	(Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-aminocarbonyl-phenyl sulphonylamino)-2-indolinone	278 (decomp.)
	ethyl	H	H	(Z)-3-(1-phenylamino-1-phenyl-methylidene)-5-ethylsulphonylamino-2-indolinone	309
	ethyl	dimethylaminomethyl	H	(Z)-3-{1-[4-dimethylaminomethyl-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone	230
	ethyl	N-benzyl-N-methyl-aminomethyl	H	(Z)-3-{1-[4-(N-benzyl-N-methyl-aminomethyl)-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone	223
	ethyl	2-dimethylamino-ethyl	H	(Z)-3-{1-[4-(2-dimethylamino-ethyl)-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone	242
	ethyl	N-(2-dimethylamino-ethyl)-N-methyl-	H	(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylsulphonyl-amino}-2-indolinone	240

					methylidene}-5-ethylsulphonylamino-2-indolinone	
ethyl		Cl		H	(Z)-3-[1-(4-chlorophenylamino)-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone	274
ethyl		nitro		H	(Z)-3-[1-[4-nitrophenylamino]-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone	270
phenyl		N-tert.butoxycarbonyl-N-ethyl-aminomethyl		H	(Z)-3-[1-[4-(N-tert.butoxycarbonyl-N-ethyl-aminomethyl)-phenylamino]-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	225
ethyl		4-(3-aminopropyl)-piperidinomethyl		H	(Z)-3-[1-[4-(4-(3-aminopropyl)-piperidinomethyl)-phenylamino]-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone	224
ethyl		4-(3-acetylamino-propyl)-piperidinomethyl		H	(Z)-3-[1-[4-(4-(3-acetylamino-propyl)-piperidinomethyl)-phenylamino]-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone	145
pyridin-3-yl		dimethylaminomethyl		H	(Z)-3-[1-[4-(4-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-(pyridin-3-ylsulphonylamino)-2-indolinone	246-247
pyridin-3-yl		pyrrolidin-1-ylmethyl		H	(Z)-3-[1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl-methylidene]-5-(pyridin-3-ylsulphonylamino)-2-indolinone	235-236

	pyridin-3-yl	N-acetyl-N-methyl-amino	H	(Z)-3-{1-[4-(N-acetyl-N-methyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	240-241
	pyridin-3-yl	N-methyl-N-methylsulphonyl-amino	H	(Z)-3-{1-[4-(N-methyl-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	286-287
	pyridin-3-yl	N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	249-250
	pyridin-3-yl	1H-imidazol-2-yl	H	(Z)-3-{1-[4-(1H-imidazol-2-yl)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	222-223
	pyridin-3-yl	1-methyl-1H-imidazol-2-yl	H	(Z)-3-{1-[4-(1-methyl-1H-imidazol-2-yl)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	230-231
	pyridin-3-yl	dimethylamino-carbonyl	H	(Z)-3-{1-[4-dimethylaminocarbonyl-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	171-172

pyridin-3-yl	4-methyl-piperazinomethyl	H	(Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	258-259
pyridin-3-yl	pyrrolidin-1-ylcarbonyl	H	(Z)-3-{1-[4-(pyrrolidin-1-ylcarbonyl)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	284-285
pyridin-3-yl	N-(2-dimethylamino-ethyl)-N-methylsulphonylamino	Cl	(Z)-3-{1-[3-chloro-4-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	261-262
pyridin-3-yl	N-(2-dimethylamino-ethyl)-N-methylsulphonylamino	NH ₂	(Z)-3-{1-[3-amino-4-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylsulphonylamino)-2-indolinone	272-273
pyridin-2-yl	4-methyl-piperazinomethyl	H	(Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	210 (decomp.)

pyridin-2-yl	N-acetyl-N-[2-(N-benzyl-N-methyl-amino)-ethyl]-amino	H	(Z)-3-{1-[4-(N-acetyl-N-(2-(N-benzyl-N-methyl-amino)-ethyl)-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	232-235
pyridin-2-yl	N-(3-dimethylamino-propyl)-N-propionyl-amino	H	(Z)-3-{1-[4-(N-(3-dimethylamino-propyl)-N-propionyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	217-219
pyridin-2-yl	N-(3-dimethylamino-propyl)-N-methylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(3-dimethylamino-propyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	258-260
pyridin-2-yl	N-(3-dimethylamino-propyl)-N-propylsulphonyl-amino	H	(Z)-3-{1-[4-(N-(3-dimethylamino-propyl)-N-propylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	256-257
pyridin-2-yl	N-(2-dimethylamino-ethyl)-N-methylsulphonyl-	H	(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	269-271

		amino]	indolinone	
pyridin-2-yl	N-piperidinomethyl-carbonyl-N-methyl-amino	H	(Z)-3-{1-[4-(N-piperidinomethylcarbonyl)-N-methyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	236-237

Example 98

(Z)-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone

5

a. 1-acetyl-3-(1-ethoxy-1-phenyl-methylidene)-5-nitro-2-indolinone

0.2 g (0.9 mmol) of 1-acetyl-5-nitro-2-indolinone and 0.6 g (2.7 mmol) of triethyl orthobenzoate are heated to 100°C in 2 ml of acetic acid anhydride for 1.5 hours. After cooling the mixture is combined with ether and the precipitate formed is suction filtered.

10 Yield: 0.2 g (66 % of theory),

Melting point: 244-250°C

R_f value: 0.7 (silica gel; ethyl acetate/cyclohexane = 3:2)

b. (Z)-1-acetyl-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-nitro-2-indolinone

15

3 g (8.5 mmol) of 1-acetyl-3-(1-ethoxy-1-phenyl-methylidene)-5-nitro-2-indolinone and 1.9 g (10 mmol) of 4-piperidinomethyl-aniline are heated to 90°C in 30 ml of DMF for 3.5 hours. After cooling to ambient temperature the reaction solution is poured onto ice water and extracted with ethyl acetate. The combined organic extracts are dried and evaporated down. The residue is triturated with ether and suction filtered.

20

Yield: 3.5 g (82 % of theory),

R_f value: 0.6 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

Melting point: 165°C

c. (Z)-1-acetyl-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-amino-2-indolinone

25

Prepared analogously to Example VIIb from (Z)-1-acetyl-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-nitro-2-indolinone by catalytic reduction over Raney nickel in dichloromethane/methanol (1:1).

30 Yield: 99 % of theory,

R_f value: 0.5 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

Melting point: 278-281°C

C₂₉H₃₀N₄O₂ (466.59)

Mass spectrum: $(M+H)^+ = 467$

d. (Z)-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone

- 5 466 mg (1 mmol) of (Z)-1-acetyl-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-amino-2-indolinone are suspended in 15 ml of pyridine, combined with 0.2 ml (2.3 mmol) of methanesulphonic acid chloride and stirred for 1.5 hours. Then 6 ml of 1 N sodium hydroxide solution are added. After 1 hour 1 ml of piperidine is added and the mixture is stirred overnight. The reaction solution is poured onto water and the
10 precipitate formed is suction filtered. The residue is stirred with ether, suction filtered and dried.

Yield: 290 mg (58 % of theory),

R_f value: 0.4 (silica gel; dichloromethane/methanol/ammonia = 9:1:0.1)

Melting point: 266°C

- 15 C₂₈H₃₀N₄O₃S (502.64)

Mass spectrum: $(M+H)^+ = 503$

Calc.: C 66.91 H 6.02 N 11.15

Found: C 66.49 H 6.06 N 11.01

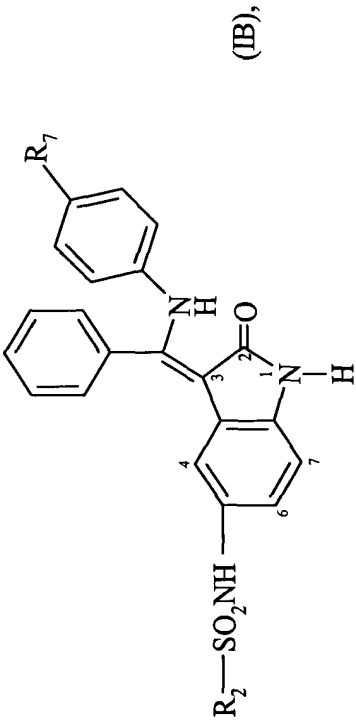
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Examples 99 to 151

- Using the intermediate products prepared in Examples I to XIII, the compounds of formula IB of Examples 99 to 151 listed in Table II are prepared analogously to Example
25 98.

Hydrochlorides or dihydrochlorides are obtained according to the following general working method: The starting compound is dissolved in dichloromethane and combined with ether/HCl. The precipitate formed is suction filtered and dried.

Table II



Example	R ₂	R ₇	chemical name	Melting point (°C)
	ethyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone	235
	ethyl	methoxy	(Z)-3-[1-(4-methoxyphenylamino)-1-phenyl-methylidene]-5-ethyl-sulphonylamino-2-indolinone	283
	isopropyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-isopropylsulphonylamino-2-indolinone	205

4-chlorophenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(4-chlorophenylsulphonylamino)-2-indolinone	251-253
3-chlorophenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-chlorophenylsulphonylamino)-2-indolinone	275-277
naphthalin-1-yl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(naphthalin-1-ylsulphonylamino)-2-indolinone	236-237
4-methylphenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(4-methylphenylsulphonylamino)-2-indolinone	267-269
3-methylphenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-methylphenylsulphonylamino)-2-indolinone	269-271
3-methoxyphenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-methoxyphenylsulphonylamino)-2-indolinone	241-245

4-methoxyphenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(4-methoxyphenylsulphonylamino)-2-indolinone	253-256
2,4,6-trimethylphenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(2,4,6-trimethylphenylsulphonylamino)-2-indolinone	224
4-nitrophenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(4-nitrophenylsulphonylamino)-2-indolinone	276
naphthalin-2-yl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(naphthalin-2-ylsulphonylamino)-2-indolinone	234
3-nitrophenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-nitrophenylsulphonylamino)-2-indolinone	145
quinolin-8-yl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(quinolin-8-ylsulphonylamino)-2-indolinone	279

2-chlorophenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(2-chlorophenylsulphonylamino)-2-indolinone	275
2-nitrophenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(2-nitrophenylsulphonylamino)-2-indolinone	140
3-cyanophenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-cyanophenylsulphonylamino)-2-indolinone	248
3,5-dimethylisoxazol-4-yl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3,5-dimethylisoxazol-4-ylsulphonylamino)-2-indolinone	240
E-2-phenylethenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-((E)-2-phenylethenylsulphonylamino)-2-indolinone	248
1-methyl-1H-imidazol-4-yl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(1-methyl-1H-imidazol-4-ylsulphonylamino)-2-indolinone-dihydrochloride	230

	cyclopropyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-cyclopropylsulphonylamino-2-indolinone	231
	2-cyanophenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(2-cyanophenylsulphonylamino)-2-indolinone	239
	pyridin-2-yl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	263
	phenyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	262
	benzyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-benzylsulphonylamino-2-indolinone	254
	propyl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-propylsulphonylamino-2-indolinone	188

benzyl	N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino	(Z)-3-{1-[4-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-phenylamino]-1-phenyl-methylidene}-5-benzylsulphonylamino-2-indolinone	163-164
isopropyl	2-dimethylamino-ethyl	(Z)-3-{1-[4-(2-dimethylamino-ethyl)-phenylamino]-1-phenyl-methylidene}-5-isopropylsulphonylamino-2-indolinone	220
propyl	2-dimethylamino-ethyl	(Z)-3-{1-[4-(2-dimethylamino-ethyl)-phenylamino]-1-phenyl-methylidene}-5-propylsulphonylamino-2-indolinone	239-240
propyl	N-benzyl-N-methyl-aminomethyl	(Z)-3-{1-[4-(N-benzyl-N-methyl-aminomethyl)-phenylamino]-1-phenyl-methylidene}-5-propylsulphonylamino-2-indolinone	195-197
methyl	N-benzyl-N-methyl-aminomethyl	(Z)-3-{1-[4-(N-benzyl-N-methyl-aminomethyl)-phenylamino]-1-phenyl-methylidene}-5-methylsulphonylamino-2-indolinone	241-242
phenyl	N-benzyl-N-methyl-aminomethyl	(Z)-3-{1-[4-(N-benzyl-N-methyl-aminomethyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	148-150

benzyl	N-benzyl-N-methyl-aminomethyl	(Z)-3-{1-[4-(N-benzyl-N-methyl-aminomethyl)-phenylamino]-1-phenyl-methylidene}-5-benzylsulphonylamino-2-indolinone	200-204
benzyl	2-dimethylamino-ethyl	(Z)-3-{1-[4-(2-dimethylamino-ethyl)-phenylamino]-1-phenyl-methylidene}-5-benzylsulphonylamino-2-indolinone-hydrochloride	260-262
pyridin-3-yl	piperidinomethyl	(Z)-3-{1-[4-(piperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(pyridin-3-ylphenylsulphonylamino)-2-indolinone	236
3-nitrophenyl	dimethylamino-methyl	(Z)-3-{1-[4-(dimethylaminomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-nitrophenylsulphonylamino)-2-indolinone	246-247
3-methoxy-phenyl	dimethylamino-methyl	(Z)-3-{1-[4-(dimethylaminomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-methoxyphenylsulphonylamino)-2-indolinone	259-260
3-nitrophenyl	dimethylamino-methyl amino	(Z)-3-{1-[4-(N-methyl-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(3-nitrophenylsulphonylamino)-2-indolinone	298-300

	2-nitrophenyl	N-methyl-N-acetyl- amino	(Z)-3-{1-[4-(N-methyl-N-acetyl-amino)-phenylamino]-1-phenyl- methylidene}-5-(2-nitrophenylsulphonylamino)-2-indolinone	295-297
	3- cyanophenyl	N-methyl-N-acetyl- amino	(Z)-3-{1-[4-(N-methyl-N-acetyl-amino)-phenylamino]-1-phenyl- methylidene}-5-(3-cyanophenylsulphonylamino)-2-indolinone	330-332
	3-nitrophenyl	4-methyl- piperazinomethyl	(Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl- methylidene}-5-(3-nitrophenylsulphonylamino)-2-indolinone	166-167
	pyridin-2-yl	pyrrolidin-1- ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl- methylidene}-5-(pyridin-2-ylsulphonylamino)-2-indolinone	261
	cyclopropyl	pyrrolidin-1- ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl- methylidene}-5-cyclopropylsulphonylamino-2-indolinone	256
	propyl	pyrrolidin-1- ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl- methylidene}-5-propylsulphonylamino-2-indolinone	247
	ethyl	pyrrolidin-1- ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl- methylidene}-5-ethylsulphonylamino-2-indolinone	245

	methyl	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenylmethylidene}-5-methylsulphonylamino-2-indolinone	248
	2-fluorophenyl	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenylmethylidene}-5-(2-fluorophenylsulphonylamino)-2-indolinone	247
	4-fluorophenyl	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenylmethylidene}-5-(4-fluorophenylsulphonylamino)-2-indolinone	244
	3-fluorophenyl	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenylmethylidene}-5-(3-fluorophenylsulphonylamino)-2-indolinone	257
	2-nitrophenyl	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenylmethylidene}-5-(2-nitrophenylsulphonylamino)-2-indolinone	185
	3-cyanophenyl	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenylmethylidene}-5-(3-cyanophenylsulphonylamino)-2-indolinone	249
	2-cyanophenyl	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenylmethylidene}-5-(2-cyanophenylsulphonylamino)-2-indolinone	232

Example 152

(Z)-3-[1-(4-ethoxycarbonylmethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone

5

a. 3-(1-ethoxy-1-phenyl-methylidene)-5-phenylsulphonylamino-2-indolinone

8 ml of 4 N sodium hydroxide solution are added to a solution of 4.0 g (8 mmol) of 1-acetyl-3-(1-ethoxy-1-phenyl-methylidene)-5-(N-acetyl-N-phenylsulphonyl-amino)-2-indolinone (Example 1e) in a mixture of 20 ml of dichloromethane and 20 ml of ethanol and the resulting mixture is stirred for 20 minutes at ambient temperature. It is then evaporated down to about. 10 ml and 150 ml of water are added. The pH is adjusted to 8-9 with 1 N hydrochloric acid. The precipitate formed is suction filtered, washed with water, isopropanol and ether, then dried *in vacuo*.

Yield: 6.6 g (82% of theory),

15 Melting point: 292-294 °C

R_f value: 0.4 (silica gel; dichloromethane/methanol/NH₄OH = 9:1:0.1)

C₂₃H₂₀N₂O₄S (420.49)

Mass spectrum: (M+H)⁺ = 421, (M-H)⁻ = 419

20 b. (Z)-3-[1-(4-ethoxycarbonylmethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone

0.84 g (2 mmol) of 3-(1-ethoxy-1-phenyl-methylidene)-5-phenylsulphonylamino-2-indolinone and 0.39 g (2.2 mmol) of 4-ethoxycarbonylmethyl-aniline are dissolved in 10 ml of DMF. The mixture is heated to 140°C for 5 hours. Then water is added while the mixture is cooled with ice and stirred for 1 hour at ambient temperature. The precipitate formed is suction filtered, washed with water, a little isopropanol and ether, then dried *in vacuo*.

Yield: 0.95 g (86 % of theory),

Melting point: 248-249°C

30 C₃₁H₂₇N₃O₅S (553.64)

Mass spectrum: M⁺ = 553, (M-H)⁻ = 552

Example 153

(Z)-3-[1-(4-carboxymethyl-phenylamino)-1-phenyl-methylidene]-5-
5 phenylsulphonylamino-2-indolinone

720 mg (1.3 mmol) of (Z)-3-[1-(4-ethoxycarbonylmethyl-phenylamino)-1-phenyl-
methylidene]-5-phenylsulphonylamino-2-indolinone are dissolved in a mixture of 20 ml
of methanol and 20 ml of dichloromethane. 4 ml of 1 N sodium hydroxide solution are
10 added and the mixture is stirred for 18 hours at ambient temperature and for another 1
hour at 40°C. The reaction solution is evaporated down to half the volume and the pH is
adjusted to 4-5 with 1 N hydrochloric acid. The precipitate formed is suction filtered,
washed with water, a little isopropanol and ether.

Yield: 620 mg (91% of theory),
15 Melting point: 305-306°C
 $C_{29}H_{23}N_3O_5S$ (525.59)
Mass spectrum: $(M-H)^+ = 524$

Example 154

20 (Z)-3-{1-[4-(benzylaminocarbonylmethyl)-phenylamino]-1-phenyl-methylidene}-5-
phenylsulphonylamino-2-indolinone

A solution of 315 mg (0.6 mmol) of (Z)-3-[1-(4-carboxymethyl-phenylamino)-1-phenyl-
25 methylidene]-5-phenylsulphonylamino-2-indolinone, 85 mg (0.8 mmol) of benzylamine,
212 mg (0.66 mmol) of TBTU and 1 ml of N-ethyl-N,N-diisopropyl-amine in 5 ml of
DMF is stirred for 3 hours at ambient temperature. Then 50 ml of water are added. The
yellow precipitate formed is suction filtered, washed with water, a little isopropanol and
ether, then dried *in vacuo*.

30 Yield: 0.3 mg (81 % of theory),
Melting point: 219-220°C

$C_{36}H_{30}N_4O_4S$ (614.73)

Mass spectrum: $(M+Na)^+ = 637$, $(M-H)^- = 613$

Example 155

5

(Z)-3-{1-[4-(N-(aminocarbonylmethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone

10 A solution of 250 mg (0.4 mmol) of (Z)-3-[1-(4-(N-carboxymethyl-N-methylsulphonyl-amino)-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone and 82 mg (0.4 mmol) of CDI in 5 ml of DMF is stirred for 1 hour at 50°C. 1 ml of condensed ammonia is added and the mixture is stirred for 5 hours at ambient temperature. Then water is added. The yellow precipitate is suction filtered, washed with water, a little isopropanol and ether, then dried *in vacuo*.

15 Yield: 190 mg (76 % of theory)

Melting point: 216-217°C

$C_{30}H_{27}N_5O_6S_2$ (617.71)

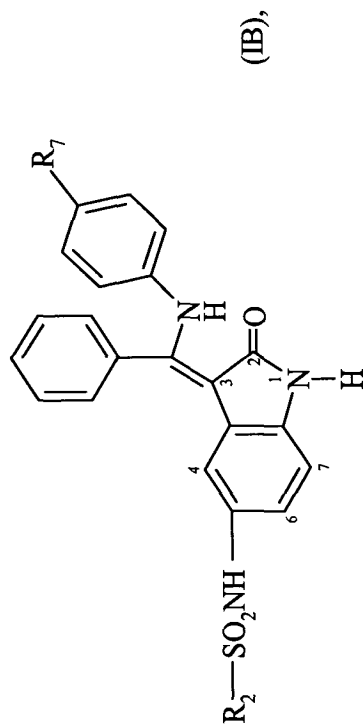
Mass spectrum: $(M+Na)^+ = 640$, $(M-H)^- = 616$

20 Examples 156 to 170

Using the intermediate products prepared in Examples I to XIII, the compounds of formula IB of Examples 156 to 170 listed in Table III are prepared analogously to Examples 152 to 155.

25

Table III



Example	R ₂	R ₇	chemical name	Melting point (°C)
	phenyl	methoxycarbonyl	(Z)-3-[1-(4-methoxycarbonyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	304-305
	phenyl	carboxy	(Z)-3-[1-(4-carboxyphenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	312-313
	phenyl	benzylaminocarbonyl	(Z)-3-{1-[4-(benzylaminocarbonyl)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	269-270
	methyl	methoxycarbonyl	(Z)-3-[1-(4-methoxycarbonyl-phenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone	> 270
	methyl	carboxy	(Z)-3-[1-(4-carboxyphenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone	> 270

phenyl	N-carboxymethyl-N-acetyl-amino	(Z)-3-{1-[4-(N-carboxymethyl-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	190-191
phenyl	N-aminocarbonylmethyl-N-acetyl-amino	(Z)-3-{1-[4-(N-(aminocarbonylmethyl)-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	150 (decomp.)
phenyl	N-methylaminocarbonylmethyl-N-acetyl-amino	(Z)-3-{1-[4-(N-methylaminocarbonylmethyl-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	150 (decomp.)
phenyl	N-dimethylaminocarbonylmethyl-N-acetyl-amino	(Z)-3-{1-[4-(N-dimethylaminocarbonylmethyl-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	150 (decomp.)
phenyl	N-carboxymethyl-N-ethylsulphonyl-amino	(Z)-3-{1-[4-(N-carboxymethyl-N-ethylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	231-235
phenyl	N-[N-(2-dimethylamino-ethyl)-N-methyl-amino-carbonylmethyl]-N-ethylsulphonyl-amino	(Z)-3-{1-[4-(N-(N-(2-dimethylamino-ethyl)-N-methyl-aminocarbonylmethyl)-N-ethylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	147-151

	phenyl	N-[(2-dimethylamino-ethyl)-aminocarbonylmethyl]-N-ethylsulphonyl-amino	(Z)-3-{1-[4-(N-((2-dimethylamino-ethyl)-aminocarbonylmethyl)-N-ethylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	142-147
	phenyl	N-carboxylmethyl-N-methylsulphonyl-amino	(Z)-3-{1-[4-(N-carboxylmethyl-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	215-216
	phenyl	N-methylaminocarbonylmethyl-N-methylsulphonyl-amino	(Z)-3-{1-[4-(N-methylaminocarbonylmethyl-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	150 (decomp.)
	phenyl	N-dimethylamino-carbonylmethyl-N-methylsulphonyl-amino	(Z)-3-{1-[4-(N-dimethylaminocarbonylmethyl-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	150 (decomp.)

Examples 171 to 206

The compounds of formula IB of Examples 171 to 206 listed in the following Table IV are obtained from compounds of the abovementioned Examples 1 to 170 by the following
5 general methods A to E or analogously to Example 1 or 210:

A: Cleaving of tert.butoxycarbonyl:

0.6 mmol of the starting compound are dissolved in 5 ml of dichloromethane. 10 ml of ethyl acetate/HCl are added and the mixture is stirred for 2 hours at ambient temperature.
10 Then a basic pH is obtained by the addition of sodium hydroxide solution. The organic phase is washed with water, dried over sodium sulphate and the solvent is eliminated *in vacuo*. In order to prepare hydrochlorides the addition of sodium hydroxide solution is omitted. In order to prepare hydrotrifluoroacetate, trifluoroacetic acid is added to the solution of the starting compound.

15 B: Cleaving of benzyl:

1.5 mmol of the starting compound are dissolved in 20 ml of dichloromethane/methanol (1:1). 100 mg of palladium/charcoal (10%) and 1.5 ml of 1 N hydrochloric acid are added and the mixture is then hydrogenated in a hydrogen atmosphere at 50 psi. The catalyst is suction filtered and the filtrate is evaporated to dryness. The residue is chromatographed
20 on silica gel (dichloromethane/methanol/NH₄OH, 9:1:0.1).

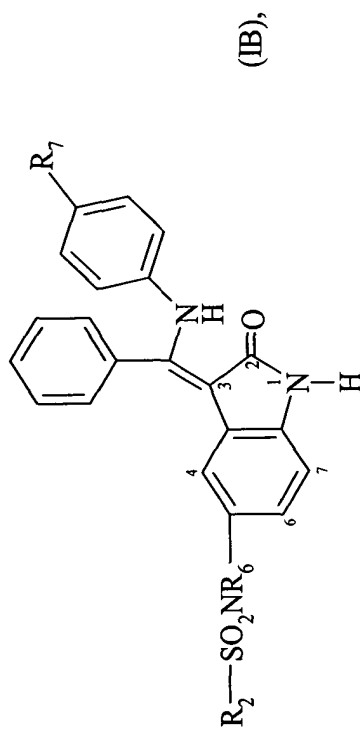
C: hydrogenation of cyano to CH₂NH₂:

0.5 mmol of the starting compound are dissolved in 20 ml of methanolic ammonia solution and combined with Raney nickel. The mixture is hydrogenated in a hydrogen
25 atmosphere of 50 psi, then the catalyst is suction filtered and the solvent is eliminated *in vacuo*. The residue is chromatographed on silica gel (dichloromethane/methanol/NH₄OH, 9:1:0.1).

D: hydrogenation of nitro to amino:

0.2 mmol of the starting compound are dissolved in 20 ml of ethyl acetate/methanol (1:1).
30 Then the mixture is hydrogenated analogously to Method C over Raney nickel. The residue is optionally chromatographed on silica gel (dichloromethane/methanol/NH₄OH, 9:1:0.1).

Table IV



Example	method	R ₂	R ₆	R ₇	chemical name	Melting point (°C)
	A	phenyl	H	amino	(Z)-3-[1-(4-aminophenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	220-223
	A	phenyl	H	piperazinomethyl	(Z)-3-[1-(4-piperazinomethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	380 (decomp.)
	A	3-methoxy-phenyl	H	aminomethyl	(Z)-3-[1-(4-aminomethyl-phenylamino)-1-phenyl-methylidene]-5-(3-methoxyphenylsulphonylamino)-2-indolinone-hydrochloride	200 (decomp.)
	A	benzyl	H	aminomethyl	(Z)-3-[1-(4-aminomethyl-phenylamino)-1-phenyl-methylidene]-5-benzylsulphonylamino)-2-indolinone-hydrochloride	200 (decomp.)

A	3-nitrophenyl	H	aminomethyl	(Z)-3-[1-(4-aminomethyl-phenylamino)-1-phenyl-methylidene]-5-(3-nitrophenylsulphonylamino)-2-indolinone-hydrochloride	215 (decomp.)
A	phenyl	H	ethylaminomethyl	(Z)-3-[1-(4-ethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone-hydrotrifluoroacetate	230
A	phenyl	H	propylamino-methyl	(Z)-3-[1-(4-propylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone-hydrotrifluoroacetate	238
A	phenyl	H	butylamino-methyl	(Z)-3-[1-(4-butylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone-hydrotrifluoroacetate	260
B	phenyl	H	N-(2-methylamino-ethyl)-N-acetyl-amino	(Z)-3-{1-[4-(N-(2-methylamino-ethyl)-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	180 (decomp.)

B	phenyl	H	N-(2-methylamino-ethyl)-N-propionyl-amino	(Z)-3-{1-[4-(N-(2-methylamino-ethyl)-N-propionyl-amino)-phenylamino]-1-phenyl-methylidene}-5-phenylsulphonylamino-2-indolinone	214
C	3-aminomethyl phenyl	H	piperidinomethyl	(Z)-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-(3-aminomethyl-phenylsulphonylamino)-2-indolinone	237
C	phenyl	H	aminomethyl	(Z)-3-[1-(4-aminomethyl-phenylamino)-1-phenyl-methylidene]-5-phenylsulphonylamino-2-indolinone	230-232
C	2-aminomethyl phenyl	H	piperidinomethyl	(Z)-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-(2-aminomethyl-phenylsulphonylamino)-2-indolinone	237
C	3-aminomethyl phenyl	H	N-methyl-N-acetyl-amino	(Z)-3-{1-[4-(N-methyl-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(3-aminomethyl-phenylsulphonylamino)-2-indolinone	277-279
C	3-aminomethyl phenyl	H	pyrrolidin-1-ylmethyl	(Z)-3-[1-(4-pyrrolidin-1-ylmethyl-phenylamino)-1-phenyl-methylidene]-5-(3-aminomethyl-phenylsulphonylamino)-2-indolinone	261

D	4-aminophenyl	H	piperidinomethyl	(Z)-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-(4-aminophenylsulphonylamino)-2-indolinone	279
D	3-aminophenyl	H	piperidinomethyl	(Z)-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-(3-aminophenylsulphonylamino)-2-indolinone	240
D	2-aminophenyl	H	piperidinomethyl	(Z)-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-(2-aminophenylsulphonylamino)-2-indolinone-hydrochloride	220 (decomp.)
D	3-aminophenyl	H	dimethylaminomethyl	(Z)-3-[1-(4-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-(3-aminophenylsulphonylamino)-2-indolinone	250 (decomp.)
D	3-aminophenyl	H	N-methyl-N-acetyl-amino	(Z)-3-{1-[4-(N-methyl-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(3-aminophenylsulphonylamino)-2-indolinone	207-209
D	2-aminophenyl	H	N-methyl-N-acetyl-amino	(Z)-3-{1-[4-(N-methyl-N-acetyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(2-aminophenylsulphonylamino)-2-indolinone	295-298

D	3-aminophenyl	H	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-aminophenylsulphonylamino)-2-indolinone	242
D	3-aminophenyl	H	(2,6-dimethylpiperidino)-methyl	(Z)-3-{1-[4-((2,6-dimethylpiperidino)-methyl)-phenylamino]-1-phenyl-methylidene}-5-(3-aminophenylsulphonylamino)-2-indolinone	150 (decomp.)
D	3-aminophenyl	H	aminomethyl	(Z)-3-[1-(4-aminomethyl-phenylamino)-1-phenyl-methylidene]-5-(3-aminophenylsulphonylamino)-2-indolinone	257
D	3-aminophenyl	H	4-methylpiperazinomethyl	(Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-(3-aminophenylsulphonylamino)-2-indolinone	217-218
D	2-aminophenyl	H	pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-(pyrrolidin-1-ylmethyl)-phenylamino]-1-phenyl-methylidene}-5-(2-aminophenylsulphonylamino)-2-indolinone	260
Ex. 1	methyl	H	acetylamino	(Z)-3-{1-[4-acetylamino-phenylamino]-1-phenyl-methylidene}-5-methylsulphonylamino-2-indolinone	299-303

Ex. 1	ethyl	H	2-dimethylamino -acetyl amino	(Z)-3-{1-[4-(2-dimethylamino-acetyl amino)-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone	238-241
Ex. 1	methyl	H	dimethylamino methyl	(Z)-3-{1-[4-dimethylaminomethyl-phenylamino]-1-phenyl-methylidene}-5-methylsulphonylamino-2-indolinone	240-242
Ex. 1	<i>n</i> -propyl	H	dimethylamino methyl	(Z)-3-{1-[4-dimethylaminomethyl-phenylamino]-1-phenyl-methylidene}-5- <i>n</i> -propylsulphonylamino-2-indolinone	221-223
Ex. 1	<i>n</i> -butyl	H	dimethylamino -methyl	(Z)-3-{1-[4-dimethylaminomethyl-phenylamino]-1-phenyl-methylidene}-5- <i>n</i> -butylsulphonylamino-2-indolinone	210-213
Ex. 1	ethyl	H	diethylamino- methyl	(Z)-3-{1-[4-diethylaminomethyl-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone	182-185
Ex. 1	ethyl	H	N-(2-dimethyl- aminoethyl)-N- methylamino- carbonyl	(Z)-3-{1-[4-(N-(2-dimethylaminoethyl)-N-methylaminocarbonyl)-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone	201-203
Ex. 210	ethyl	CH ₃	dimethylamino -methyl	(Z)-3-[1-(4-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-(N-ethyl-N-phenylsulphonyl-amino)-2-indolinone	

Ex. 1	pyrid-2-yl	H	(S)-2-hydroxy-methyl-pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-((S)-2-hydroxymethylpyrrolid-1-ylmethyl)-phenylamino]-1-phenyl-methylidene}-5-pyrid-2-ylsulphonylamino-2-indolinone	sintering from 100
Ex. 1	pyrid-2-yl	H	(S)-3-hydroxy-pyrrolidin-1-ylmethyl	(Z)-3-{1-[4-((S)-3-hydroxypyrrolid-1-ylmethyl)-phenylamino]-1-phenyl-methylidene}-5-pyrid-2-ylsulphonylamino-2-indolinone	sintering from 130

Example 207

(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(N-methyl-N-phenylsulphonyl-amino)-2-indolinone

5

a. (Z)-1-acetyl-3-(1-ethoxy-1-phenyl-methylidene)-5-(N-methyl-N-phenylsulphonyl-amino)-2-indolinone

10 g (20 mmol) of 1-acetyl-3-(1-ethoxy-1-phenyl-methylidene)-5-(N-acetyl-N-phenylsulphonyl-amino)-2-indolinone (Example 1e) are dissolved in 150 ml of DMSO and combined with 2.2 g (20 mmol) of potassium tert. butoxide with stirring. After 15 minutes' stirring 1.9 ml (31 mmol) of iodomethane are added. The mixture is stirred for 3 hours at ambient temperature. Then another 2.2 g (20 mmol) of potassium tert. butoxide and 1 ml (16 mmol) of iodomethane are added. The mixture is stirred for 18 hours at ambient temperature. Then water is added. The reaction mixture is extracted with ethyl acetate. The organic phase is washed with water, dried over magnesium sulphate and evaporated to dryness. The residue is chromatographed on silica gel (petroleum ether/dichloromethane, 7:3).

15

Yield: 2.7 g (28% of theory)

R_f value: 0.65 (silica gel; dichloromethane/petroleum ether = 8:2)

20

C₂₆H₂₄N₂O₅S (476.56)

Mass spectrum: (M+Na)⁺ = 499

b. (Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino)-phenylamino]-1-phenyl-methylidene}-5-(N-methyl-N-phenylsulphonyl-amino)-2-indolinone

25

Prepared analogously to Example 1f from 350 mg (0.73 mmol) of (Z)-1-acetyl-3-(1-ethoxy-1-phenyl-methylidene)-5-(N-methyl-N-phenylsulphonyl-amino)-2-indolinone and 257 mg (1 mmol) of 4-[N-(2-dimethylamino-ethyl)-N-methylsulphonyl-amino]-aniline in DMF and subsequent treatment with sodium hydroxide solution.

Yield: 380 mg (80% of theory)

30

R_f value: 0.5 (silica gel; dichloromethane/methanol/NH₄OH = 9:1:0.1)

C₃₃H₃₅N₅O₅S₂ (645.80)

Mass spectrum: M⁺ = 645

Calc.: C 61.38 H 5.46 N 10.84

Found: C 61.09 H 5.45 N 10.82

The following compounds of Examples 208 to 210 are prepared analogously to Example 207 using the intermediate products prepared in Examples I to XIII:

5

Example 208

(Z)-3-{1-[4-(N-(2-dimethylamino-ethyl)-N-acetyl-amino)-phenylamino)-1-phenyl-methylidene]-5-(N-methyl-N-phenylsulphonyl-amino)-2-indolinone

10

Melting point: 217°C

R_f value: 0.5 (silica gel; dichloromethane/methanol/NH₄OH = 9:1:0.1)

C₃₄H₃₅N₅O₄S (609.75)

Mass spectrum: (M+H)⁺ = 610

15

Calc.: C 66.97 H 5.79 N 11.49

Found: C 66.92 H 5.78 N 11.39

Example 209

20

(Z)-3-{1-[4-(N-methyl-N-piperidinomethylcarbonyl-amino)-phenylamino)-1-phenyl-methylidene]-5-(N-methyl-N-phenylsulphonyl-amino)-2-indolinone

Melting point: 160°C

R_f value: 0.65 (silica gel; dichloromethane/methanol/NH₄OH = 9:1:0.1)

25

C₃₆H₃₇N₅O₄S (635.79)

Mass spectrum: (M+H)⁺ = 636

Example 210

30

(Z)-3-[1-(3-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-(N-methyl-N-phenylsulphonyl-amino)-2-indolinone

Melting point: 226°C

R_f value: 0.75 (silica gel; dichloromethane/methanol/NH₄OH = 9:1:0.1)

C₃₁H₃₀N₄O₃S (538.67)

Mass spectrum: (M+H)⁺ = 539

5 The following compounds may be obtained analogously to the foregoing Examples:

- (1) (Z)-3-[1-(3-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone, melting point 222-224 °C
- (2) (Z)-3-{1-[4-(2-dimethylaminoethyl)-phenylamino]-1-phenyl-methylidene}-5-methylsulphonylamino-2-indolinone
- 10 (3) (Z)-3-[1-(4-diethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone
- (4) (Z)-3-[1-(4-dipropylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone
- 15 (5) (Z)-3-[1-(4-hexamethyleneiminomethyl-phenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone
- (6) (Z)-3-{1-[4-(4-methylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-methylsulphonylamino-2-indolinone
- (7) (Z)-3-[1-(4-morpholinomethyl-phenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone
- 20 (8) (Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-methylsulphonylamino-2-indolinone
- (9) (Z)-3-[1-(4-piperazinomethyl-phenylamino)-1-phenyl-methylidene]-5-methylsulphonylamino-2-indolinone
- 25 (10) (Z)-3-{1-[4-(2,6-dimethylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-methylsulphonylamino-2-indolinone
- (11) (Z)-3-[1-(4-dipropylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone
- (12) (Z)-3-[1-(4-hexamethyleneiminomethyl-phenylamino)-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone
- 30 (13) (Z)-3-{1-[4-(4-methylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone

- (14) (Z)-3-[1-(4-morpholinomethyl-phenylamino)-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone
- (15) (Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone
- 5 (16) (Z)-3-[1-(4-piperazinomethyl-phenylamino)-1-phenyl-methylidene]-5-ethylsulphonylamino-2-indolinone
- (17) (Z)-3-{1-[4-(2,6-dimethylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-ethylsulphonylamino-2-indolinone
- (18) (Z)-3-[1-(4-piperazinomethyl-phenylamino)-1-phenyl-methylidene]-5-propylsulphonylamino-2-indolinone
- 10 (19) (Z)-3-{1-[4-(2,6-dimethylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-propylsulphonylamino-2-indolinone
- (20) (Z)-3-[1-(4-diethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-propylsulphonylamino-2-indolinone
- 15 (21) (Z)-3-[1-(4-dipropylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-propylsulphonylamino-2-indolinone
- (22) (Z)-3-[1-(4-hexamethyleneiminomethyl-phenylamino)-1-phenyl-methylidene]-5-propylsulphonylamino-2-indolinone
- (23) (Z)-3-{1-[4-(4-methylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-propylsulphonylamino-2-indolinone
- 20 (24) (Z)-3-[1-(4-morpholinomethyl-phenylamino)-1-phenyl-methylidene]-5-propylsulphonylamino-2-indolinone
- (25) (Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-propylsulphonylamino-2-indolinone
- 25 (26) (Z)-3-[1-(4-piperazinomethyl-phenylamino)-1-phenyl-methylidene]-5-cyclopropylsulphonylamino-2-indolinone
- (27) (Z)-3-{1-[4-(2,6-dimethylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-cyclopropylsulphonylamino-2-indolinone
- (28) (Z)-3-[1-(4-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-cyclopropylsulphonylamino-2-indolinone
- 30 (29) (Z)-3-{1-[4-(2-dimethylaminoethyl)-phenylamino]-1-phenyl-methylidene}-5-cyclopropylsulphonylamino-2-indolinone

- (30) (Z)-3-[1-(4-diethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-cyclopropylsulphonylamino-2-indolinone
- (31) (Z)-3-[1-(4-dipropylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-cyclopropylsulphonylamino-2-indolinone
- 5 (32) (Z)-3-[1-(4-hexamethyleneiminomethyl-phenylamino)-1-phenyl-methylidene]-5-cyclopropylsulphonylamino-2-indolinone
- (33) (Z)-3-{1-[4-(4-methylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-cyclopropylsulphonylamino-2-indolinone
- (34) (Z)-3-[1-(4-morpholinomethyl-phenylamino)-1-phenyl-methylidene]-5-cyclopropylsulphonylamino-2-indolinone
- 10 (35) (Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-cyclopropylsulphonylamino-2-indolinone
- (36) (Z)-3-[1-(4-piperazinomethyl-phenylamino)-1-phenyl-methylidene]-5-trifluoromethylsulphonylamino-2-indolinone
- 15 (37) (Z)-3-{1-[4-(2,6-dimethylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-trifluoromethylsulphonylamino-2-indolinone
- (38) (Z)-3-[1-(4-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-trifluoromethylsulphonylamino-2-indolinone
- (39) (Z)-3-{1-[4-(2-dimethylaminoethyl)-phenylamino]-1-phenyl-methylidene}-5-trifluoromethylsulphonylamino-2-indolinone
- 20 (40) (Z)-3-[1-(4-diethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-trifluoromethylsulphonylamino-2-indolinone
- (41) (Z)-3-[1-(4-dipropylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-trifluoromethylsulphonylamino-2-indolinone
- 25 (42) (Z)-3-{1-[4-(pyrrolidin-1-yl)-methyl-phenylamino]-1-phenyl-methylidene}-5-trifluoromethylsulphonylamino-2-indolinone
- (43) (Z)-3-[1-(4-piperidinomethyl-phenylamino)-1-phenyl-methylidene]-5-trifluoromethylsulphonylamino-2-indolinone
- (44) (Z)-3-[1-(4-hexamethyleneiminomethyl-phenylamino)-1-phenyl-methylidene]-5-trifluoromethylsulphonylamino-2-indolinone
- 30 (45) (Z)-3-{1-[4-(4-methylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-trifluoromethylsulphonylamino-2-indolinone

- (46) (Z)-3-[1-(4-morpholinomethyl-phenylamino)-1-phenyl-methylidene]-5-trifluoromethylsulphonylamino-2-indolinone
- (47) (Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-trifluoromethylsulphonylamino-2-indolinone
- 5 (48) (Z)-3-[1-(4-piperazinomethyl-phenylamino)-1-phenyl-methylidene]-5-isopropylsulphonylamino-2-indolinone
- (49) (Z)-3-{1-[4-(2,6-dimethylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-isopropylsulphonylamino-2-indolinone
- (50) (Z)-3-[1-(4-dimethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-isopropylsulphonylamino-2-indolinone
- 10 (51) (Z)-3-[1-(4-diethylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-isopropylsulphonylamino-2-indolinone
- (52) (Z)-3-[1-(4-dipropylaminomethyl-phenylamino)-1-phenyl-methylidene]-5-isopropylsulphonylamino-2-indolinone
- 15 (53) (Z)-3-{1-[4-(pyrrolidin-1-yl)-methyl-phenylamino]-1-phenyl-methylidene}-5-isopropylsulphonylamino-2-indolinone
- (54) (Z)-3-[1-(4-hexamethyleneiminomethyl-phenylamino)-1-phenyl-methylidene]-5-isopropylsulphonylamino-2-indolinone
- (55) (Z)-3-{1-[4-(4-methylpiperidinomethyl)-phenylamino]-1-phenyl-methylidene}-5-isopropylsulphonylamino-2-indolinone
- 20 (56) (Z)-3-[1-(4-morpholinomethyl-phenylamino)-1-phenyl-methylidene]-5-isopropylsulphonylamino-2-indolinone
- (57) (Z)-3-{1-[4-(4-methylpiperazinomethyl)-phenylamino]-1-phenyl-methylidene}-5-isopropylsulphonylamino-2-indolinone
- 25

Example 211

Dry ampoule containing 75 mg of active substance per 10 ml

30 Composition:

Active substance	75.0 mg
Mannitol	50.0 mg

water for injections ad 10.0 ml

Preparation:

Active substance and mannitol are dissolved in water. After packaging the solution is
5 freeze-dried. To produce the solution ready for use, the product is dissolved in water for
injections.

Example 212

10 Dry ampoule containing 35 mg of active substance per 2 ml

Composition:

Active substance	35.0 mg
Mannitol	100.0 mg
15 water for injections	ad 2.0 ml

Preparation:

Active substance and mannitol are dissolved in water. After packaging, the solution is
freeze-dried.

20

To produce the solution ready for use, the product is dissolved in water for injections.

Example 213

25

Tablet containing 50 mg of active substance

Composition:

30 (1) Active substance	50.0 mg
(2) Lactose	98.0 mg
(3) Maize starch	50.0 mg
(4) Polyvinylpyrrolidone	15.0 mg

(5) Magnesium stearate 2.0 mg
 215.0 mg

Preparation:

- 5 (1), (2) and (3) are mixed together and granulated with an aqueous solution of (4). (5) is added to the dried granulated material. From this mixture tablets are pressed, biplanar, faceted on both sides and with a dividing notch on one side.
Diameter of the tablets: 9 mm.

10

Example 214

Tablet containing 350 mg of active substance

- 15 Preparation:

(1) Active substance 350.0 mg
(2) Lactose 136.0 mg
(3) Maize starch 80.0 mg
20 (4) Polyvinylpyrrolidone 30.0 mg
(5) Magnesium stearate 4.0 mg
 600.0 mg

- (1), (2) and (3) are mixed together and granulated with an aqueous solution of (4). (5) is
25 added to the dried granulated material. From this mixture tablets are pressed, biplanar, faceted on both sides and with a dividing notch on one side.
Diameter of the tablets: 12 mm.

Example 215

30

Capsules containing 50 mg of active substance

Composition:

	(1) Active substance	50.0 mg
	(2) Dried maize starch	58.0 mg
	(3) Powdered lactose	50.0 mg
	(4) Magnesium stearate	2.0 mg
5		160.0 mg

Preparation:

(1) is triturated with (3). This trituration is added to the mixture of (2) and (4) with vigorous mixing.

10

This powder mixture is packed into size 3 hard gelatine capsules in a capsule filling machine.

Example 216

15

Capsules containing 350 mg of active substance

Composition:

	(1) Active substance	350.0 mg
20	(2) Dried maize starch	46.0 mg
	(3) Powdered lactose	30.0 mg
	(4) Magnesium stearate	4.0 mg
		430.0 mg

25 Preparation:

(1) is triturated with (3). This trituration is added to the mixture of (2) and (4) with vigorous mixing.

This powder mixture is packed into size 0 hard gelatine capsules in a capsule filling machine.

30

Example 217

Suppositories containing 100 mg of active substance

1 suppository contains:

	active substance	100.0 mg
5	polyethyleneglycol (M.W. 1500)	600.0 mg
	polyethyleneglycol (M.W. 6000)	460.0 mg
	polyethylenesorbitan monostearate	840.0 mg
		2,000.0 mg

10 Preparation:

The polyethyleneglycol is melted together with polyethylene sorbitan monostearate. At 40°C the ground active substance is homogeneously dispersed in the melt. It is cooled to 38°C and poured into slightly chilled suppository moulds.